Print selected from Online session 22/03/2004Page 1

=> d hist

(FILE 'HOME' ENTERED AT 18:25:05 ON 22 MAR 2004)

FILE 'REGISTRY' ENTERED AT 18:25:18 ON 22 MAR 2004 E LITHIUM IRON MAGNESIUM PHOSPHATE/CN E IRON LITHIUM MAGNESIUM PHOSPHATE/CN

L1 22 S E3-24

FILE 'CAPLUS' ENTERED AT 18:27:25 ON 22 MAR 2004 L2 30 S L1

=> d ibib ab it 1-

YOU HAVE REQUESTED DATA FROM 30 ANSWERS - CONTINUE? Y/(N):y

L2 ANSWER 1 OF 30 CAPLUS COPYRIGHT 2004 ACS on STN

ACCESSION NUMBER:

2004:78230 CAPLUS

DOCUMENT NUMBER:

140:114274

TITLE:

0

Methods of making transition metal compounds useful as

battery cathode active materials using electromagnetic

radiation

INVENTOR(S):

Barker, Jeremy

PATENT ASSIGNEE(S):

UK

SOURCE:

U.S. Pat. Appl. Publ., 23 pp.

CODEN: USXXCO

DOCUMENT TYPE:

Patent

LANGUAGE:

English

FAMILY ACC. NUM. COUNT: 1

PATENT INFORMATION:

PATENT NO. KIND DATE APPLICATION NO. DATE

US 2004016632 A1 20040129 US 2002-205748 20020726

PRIORITY APPLN. INFO.: US 2002-205748 20020726

AB In a method for synthesizing reduced metal compds. using electromagnetic radiation, starting materials comprising at least one particulate metal compound and at least one source of carbon are combined to form a mixture. The mixture is exposed to electromagnetic radiation to form a reaction product. Preferably, the carbon is a reducing carbon, and at least one metal of the starting materials is reduced in oxidation state during radiation exposure. Reducing carbon may be supplied by elemental carbon, by an organic material, or by mixts. Preferably, the solid state reactants also include an alkali metal compound. The products of the method are preferably useful as cathode active materials in lithium ion batteries. The electromagnetic radiation is selected from among microwave, IR, and radio frequencies of .apprx.1 MHz to 3000 GHz.

IT Secondary batteries

(lithium; methods of making transition metal compds. useful as battery cathode active materials using electromagnetic radiation)

IT Battery cathodes

Electromagnetic wave

```
Solid phase synthesis
       (methods of making transition metal compds. useful as battery cathode
       active materials using electromagnetic radiation)
    Alkali metal compounds
IT
    Carbohydrates, processes
    Carbon black, processes
    RL: CPS (Chemical process); PEP (Physical, engineering or chemical
    process); PROC (Process)
        (methods of making transition metal compds, useful as battery cathode
       active materials using electromagnetic radiation)
    Transition metal compounds
IT
    RL: DEV (Device component use); SPN (Synthetic preparation); PREP
    (Preparation): USES (Uses)
        (methods of making transition metal compds. useful as battery cathode
       active materials using electromagnetic radiation)
    Transition metal oxides
IT
    RL: DEV (Device component use); SPN (Synthetic preparation); PREP
     (Preparation): USES (Uses)
        (methods of making transition metal compds. useful as battery cathode
        active materials using electromagnetic radiation)
ΙT
    Materials
        (organic; methods of making transition metal compds. useful as battery
        cathode active materials using electromagnetic radiation)
    57-50-1, Sucrose, processes 7439-89-6D, Iron, compound 7439-93-2D,
ΙT
    Lithium, compound 7439-96-5D, Manganese, compound 7439-98-7D, Molybdenum,
               7440-02-0D, Nickel, compound 7440-31-5D, Tin, compound
    7440-32-6D. Titanium, compound
                                     7440-44-0, Carbon, processes
                                                                    7440-47-3D.
                        7440-48-4D, Cobalt, compound
                                                       7440-50-8D. Copper. compound
    Chromium, compound
    7440-62-2D. Vanadium, compound 7782-42-5. Graphite, processes
    RL: CPS (Chemical process); PEP (Physical, engineering or chemical
    process); PROC (Process)
        (methods of making transition metal compds. useful as battery cathode
        active materials using electromagnetic radiation)
    11126-15-1, Lithium vanadium oxide 37296-91-6, Lithium molybdenum oxide
ΙT
     39302-37-9, Lithium titanium oxide
    RL: DEV (Device component use); USES (Uses)
        (methods of making transition metal compds. useful as battery cathode
        active materials using electromagnetic radiation)
    7664-38-2DP. Phosphoric acid. transition metal compds.
IT
    Lithium molybdenum oxide limoo2 84159-18-2P, Lithium vanadium phosphate
    Li3V2(PO4)3 160014-96-OP, Lithium molybdenum oxide Li4Mo308
     610271-94-8P 632286-77-2P. Iron lithium magnesium phosphate
     (Fe0.9LiMg0.1P04)
     RL: DEV (Device component use); SPN (Synthetic preparation); PREP
     (Preparation); USES (Uses)
        (methods of making transition metal compds. useful as battery cathode
        active materials using electromagnetic radiation)
    124-38-9, Carbon dioxide, formation (nonpreparative) 630-08-0, Carbon
IT
    monoxide, formation (nonpreparative)
```

```
RL: FMU (Formation, unclassified); FORM (Formation, nonpreparative) (methods of making transition metal compds. useful as battery cathode active materials using electromagnetic radiation)
1333-74-0, Hydrogen, processes
RL: CPS (Chemical process); PEP (Physical, engineering or chemical process); PROC (Process)
(reducing gas; methods of making transition metal compds. useful as battery cathode active materials using electromagnetic radiation)
12162-92-4P. Lithium vanadiumoxide LiV205
RL: DEV (Device component use); SPN (Synthetic preparation); PREP (Preparation); USES (Uses)
```

 $(\gamma$ -; methods of making transition metal compds. useful as battery

_2 ANSWER 2 OF 30 CAPLUS COPYRIGHT 2004 ACS on STN

ACCESSION NUMBER:

2004:59636 CAPLUS

cathode active materials using electromagnetic radiation)

DOCUMENT NUMBER:

140:96916

TITLE:

Method of synthesizing electrochemically active

materials from a slurry of precursors

INVENTOR(S):

Stoker, John; Hodge, James

PATENT ASSIGNEE(S):

USA

SOURCE:

U.S. Pat. Appl. Publ., 17 pp.

CODEN: USXXCO

DOCUMENT TYPE:

Patent

LANGUAGE:

English

FAMILY ACC. NUM. COUNT: 1

PATENT INFORMATION:

PATENT NO.	KIND DAT	E	APPLICATION	N NO. DA	ATE			
US 2004013943 WO 2004010515			US 2002-200		•			
W: AE, AG,	AL, AM, AT	, AU, AZ,	BA, BB, BG, I DZ, EC, EE, I	BR, BY, E	BZ, CA, CH,			
GM, HR,	HU, ID, IL	, IN, IS,	JP, KE, KG, I	KP, KR, k	KZ, LC, LK,	LR,		
			MK, MN, MW, 1 SD, SE, SG, S					
TR, TT, KG, KZ,		, US, UZ,	VC, VN, YU, 7	ZA, ZM, Z	ZW, AM, AZ,	BY,		
RW: GH, GM,	KE, LS, MW		SL, SZ, TZ, U FI, FR, GB, (
NL, PT.	RO, SE, SI	, SK, TR,	BF, BJ, CF, (
	GW, ML. MR, NE, SN, TD, TG RIORITY APPLN. INFO.: US 2002-200823 A 20020722							
AB A method for mal	Ū			•		_		

AB A method for making an active material comprises the steps of forming a slurry, spray drying the slurry to form a powdered precursor composition, and heating the powdered precursor composition at a temperature and for a time sufficient to

form a reaction product. The slurry has a liquid phase and a solid phase, and contains at least an alkali metal compound and a transition metal compound

IT

ΙT

ΙT

ΙT

IT

precursors)

Preferably the liquid phase contains dissolved alkali metal compound, and the solid phase contains an insol. transition metal compound, an insol. carbonaceous material compound, or both. Electrodes and batteries are provided that contain the active materials. Dispersing agents (anionic; method of synthesizing electrochem. active materials from slurry of precursors) Battery electrodes Secondary batteries Slurries (method of synthesizing electrochem. active materials from slurry of precursors) Alkali metal compounds Carbohydrates, uses Carbonaceous materials (technological products) Hydrocarbons, uses Polymers, uses Transition metal compounds RL: DEV (Device component use); USES (Uses) (method of synthesizing electrochem. active materials from slurry of precursors) 25155-19-5D, Naphthalene sulfonic acid, salt RL: MOA (Modifier or additive use); USES (Uses) (dispersant: method of synthesizing electrochem. active materials from slurry of precursors) 7439-88-5D, Iridium, compound 7439-89-6D, Iron, compound 7439-93-2D, Lithium. compound 7439-95-4D. Magnesium, compound 7439-96-5D. Manganese. compound 7439-98-7D, Molybdenum, compound 7440-02-0D, Nickel, compound 7440-04-2D. Osmium, compound 7440-15-5D, Rhenium, compound 7440-24-6D. Strontium, compound 7440-33-7D, Tungsten, compound 7440-39-3D, Barium, compound 7440-41-7D, Beryllium, compound 7440-44-0, Carbon, uses 7440-47-3D, Chromium, compound 7440-48-4D, Cobalt, compound 7440-62-2D. Vanadium, compound 7440-66-6D, Zinc, compound 7440-70-2D, Calcium, compound 14066-19-4. Hydrogen Phosphate, uses 14066-20-7. DiHydrogen Phosphate. 14265-44-2, Phosphate, uses 15365-14-7, Iron lithium phosphate uses felipo4 RL: DEV (Device component use); USES (Uses) (method of synthesizing electrochem. active materials from slurry of precursors) 349632-85-5P 632286-77-2P, Iron lithium magnesium phosphate Fe0.9LiMg0.1P04 643752-34-5P. Iron lithium magnesium phosphate (Fe0.95LiMg0.05(PO4)) RL: DEV (Device component use); SPN (Synthetic preparation); PREP (Preparation): USES (Uses) (method of synthesizing electrochem. active materials from slurry of

25852-26-0D. Formaldehyde-naphthalene copolymer, sodium sulfonate 25852-26-0D. Naphthalene formaldehyde copolymer, sulfonic acid salts RL: MOA (Modifier or additive use); USES (Uses) (method of synthesizing electrochem. active materials from slurry of

precursors)

IT 7732-18-5, Water, uses

RL: TEM (Technical or engineered material use); USES (Uses) (method of synthesizing electrochem. active materials from slurry of precursors)

L2 ANSWER 3 OF 30 CAPLUS COPYRIGHT 2004 ACS on STN

ACCESSION NUMBER:

2003:950942 CAPLUS

DOCUMENT NUMBER:

139:398062

TITLE:

Synthesis of metal compounds useful as cathode active

materials

INVENTOR(S):

Barker, Jeremy; Saidi, Yazid M.; Swoyer, Jeffrey:

Dong, Ming

PATENT ASSIGNEE(S):

Valence Technology, Inc., USA

SOURCE:

PCT Int. Appl., 78 pp.

CODEN: PIXXD2

DOCUMENT TYPE:

Patent

LANGUAGE:

English

FAMILY ACC. NUM. COUNT: 2

PATENT INFORMATION:

```
APPLICATION NO.
    PATENT NO.
                     KIND
                           DATE
                                                            DATE
                                           WO 2003-US14290 20030506
                            20031204
    WO 2003099715
                      A1
        W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN,
            CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, ES, FI, GB, GD, GE, GH,
            GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR.
            LS. LT. LU. LV. MA. MD. MG. MK, MN, MW, MX, MZ, NO, NZ, OM, PH.
            PL, PT, RO, RU, SC, SD, SE, SG, SK, SL, TJ, TM, TN, TR, TT, TZ,
            UA, UG, US, UZ, VC, VN, YU, ZA, ZM, ZW, AM, AZ, BY, KG, KZ, MD,
            RU. TJ, TM
         RW: GH. GM. KE. LS. MW. MZ, SD, SL, SZ, TZ, UG, ZM, ZW, AT, BE, BG,
            CH. CY. CZ. DE. DK. EE. ES. FI. FR. GB, GR. HU, IE, IT, LU, MC.
            NL, PT, RO, SE, SI, SK, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ,
            GW, ML, MR, NE, SN, TD, TG
                                           US 2002-150353
                                                            20020517
     US 2003215715
                      A1
                            20031120
PRIORITY APPLN. INFO.:
                                        US 2002-150343
                                                        A 20020517
```

Active materials of the invention contain at least one alkali metal and at least one other metal capable of being oxidized to a higher oxidation state. Preferred other metals are accordingly selected from the group consisting of transition metals (defined as Groups 4-11 of the periodic table), as well as certain other nontransition materials such as tin, bismuth, and lead. The active materials may be synthesized in single step reactions or in multi-step reactions. In at least one of the steps of the synthesis reaction, reducing carbon is used as a starting material. In one aspect, the reducing carbon is provided by elemental carbon, preferably in particulate form such as graphites, amorphous carbon, carbon blacks and the like. In another aspect, reducing carbon may also be provided by an organic precursor material, or by a mixture of elemental carbon and organic

```
precursor material.
    Reduction
IT
       (carbothermic; synthesis of metal compds. useful as cathode active
       materials)
    Oxides (inorganic), preparation
    Phosphates, preparation
    RL: SPN (Synthetic preparation): PREP (Preparation)
       (lithiated; synthesis of metal compds. useful as cathode active
       materials)
   Battery cathodes
       (synthesis of metal compds. useful as cathode active materials)
IT Alkali metal compounds
    Butadiene rubber, processes
    Carbohydrates, processes
    Hydrocarbons, processes
    Transition metal compounds
    RL: CPS (Chemical process); PEP (Physical, engineering or chemical
    process); PROC (Process)
       (synthesis of metal compds. useful as cathode active materials)
    Carbonaceous materials (technological products)
IT
    RL: CPS (Chemical process); PEP (Physical, engineering or chemical
    process); SPN (Synthetic preparation); PREP (Preparation); PROC (Process)
       (synthesis of metal compds. useful as cathode active materials)
    9003-17-2
ΙT
    RL: CPS (Chemical process); PEP (Physical, engineering or chemical
    process); PROC (Process)
       (butadiene rubber, synthesis of metal compds. useful as cathode active
       materials)
IT
    57-50-1, Sucrose, processes 554-13-2, Lithium carbonate
    Ferric oxide, processes 1310-66-3, Lithium hydroxide monohydrate
     1313-27-5. Molybdenum oxide moo3, processes 1314-62-1, Vanadium oxide
     (V205), processes 1333-74-0, Hydrogen, processes 7439-89-6D. Iron,
              7439-93-2D. Lithium, compound 7439-96-5D, Manganese, compound
    7439-98-7D, Molybdenum, compound 7440-02-0D, Nickel, compound 7440-31-5D.
    Tin, compound 7440-32-6D, Titanium, compound 7440-44-0, Carbon, processes
     7440-47-3D. Chromium. compound 7440-48-4D. Cobalt. compound 7440-50-8D.
    Copper. compound 7440-62-2D, Vanadium, compound 14940-41-1, Ferrous
     phosphate 84159-18-2. Lithium vanadium phosphateLi3V2(PO4)3
     RL: CPS (Chemical process); PEP (Physical, engineering or chemical
     process): PROC (Process)
        (synthesis of metal compds. useful as cathode active materials)
     18868-43-4P. Molybdenum oxide (MoO2)
     RL: CPS (Chemical process); PEP (Physical, engineering or chemical
     process); SPN (Synthetic preparation); PREP (Preparation); PROC (Process)
       (synthesis of metal compds. useful as cathode active materials)
    7664-38-2DP. Phosphoric acid, transition metal compds. 11126-15-1P,
IT
     Lithium vanadium oxide 12162-92-4P, Lithium vanadium oxide liv2o5
     15365-14-7P, Iron lithium phosphate felipo4 37296-91-6P, Lithium
     molybdenum oxide 39302-37-9P. Lithium titanium oxide 39457-42-6P.
     Lithium manganese oxide 69550-44-3P, Lithium molybdenum oxide limoo2
```

108399-57-1P, Lithium molybdenum oxide Li0.74MoO2 349632-76-4P. Iron lithium magnesium phosphate (Fe0.9LiMg0.1(PO4)) 349632-79-7P. Calcium iron lithium phosphate Ca0.1Fe0.9LiPO4 507242-52-6P. Lithium molybdenum oxide LiO-2MoO2 507242-65-1P. Lithium molybdenum oxide Li0.85Mo02 RL: SPN (Synthetic preparation); PREP (Preparation)

(synthesis of metal compds. useful as cathode active materials)

REFERENCE COUNT:

4 THERE ARE 4 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

L2 ANSWER 4 OF 30 CAPLUS COPYRIGHT 2004 ACS on STN

ACCESSION NUMBER:

2003:947998 CAPLUS

DOCUMENT NUMBER:

140:29489

TITLE:

Nonaqueous electrolyte secondary battery using nickel

metal oxide as anode active mass

INVENTOR(S):

Kitano, Shinya

PATENT ASSIGNEE(S):

Japan Storage Battery Co., Ltd., Japan

SOURCE:

Jpn. Kokai Tokkyo Koho, 11 pp.

CODEN: JKXXAF

DOCUMENT TYPE:

Patent

LANGUAGE:

Japanese

FAMILY ACC. NUM. COUNT: 1

PATENT INFORMATION:

PATENT NO. APPLICATION NO. DATE KIND DATE Α2 JP 2002-151010 20020524 20031205 JP 2003346799 JP 2002-151010 20020524 PRIORITY APPLN. INFO.:

The non-aqueous electrolytic battery comprises an anode containing an anode active mass containing LiCoO2 50-80, LiMnO2 10-30, and LiFe1-xMnxPO4 (0 \leq x ≤ 0.13 ; M = Mg, Co, Ni, Mn, and/or Zn) 10-30 weight%. The non-aqueous electrolytic battery comprising the anode active mass is provided with excellent overcharging property and shows continuous voltage alteration at the time charging and discharging.

Secondary batteries ΙT

> (lithium; nonag. electrolyte batter containing lithium compound mixture as anode active mass for good overcharging property)

Battery anodes IT

> (of nonaq. electrolyte secondary battery; nonaq. electrolyte batter containing lithium compound mixture as anode active mass for good overcharging property)

12162-79-7, Lithium manganese oxide (LiMnO2) 12190-79-3, Cobalt lithium oxide (CoLiO2) 15365-14-7, Iron lithium phosphate (LiFePO4) 412351-36-1. Iron lithium manganese phosphate (Fe0.9LiMn0.1(PO4))

632286-78-3 632286-79-4 632286-80-7 632286-77-2 RL: TEM (Technical or engineered material use); USES (Uses)

(anode active mass containing; nonaq. electrolyte batter containing lithium compound mixture as anode active mass for good overcharging property)

Print selected from Online session 22/03/2004Page 8

ACCESSION NUMBER:

2003:912686 CAPLUS

DOCUMENT NUMBER:

139:367603

TITLE:

Methods of making transition metal compounds useful as

battery cathode active materials

INVENTOR(S):

Barker, Jeremy; Saidi, M. Yazid; Swoyer, Jeffrey L.;

Dong, Ming

PATENT ASSIGNEE(S):

Valence Technology, Inc., USA

SOURCE:

U.S. Pat. Appl. Publ., 22 pp.

CODEN: USXXCO

DOCUMENT TYPE:

Patent

LANGUAGE:

English

FAMILY ACC. NUM. COUNT:

PATENT INFORMATION:

PATENT NO.	KIND DATE	•	APPLICATION	NO. DATE	
US 2003215715	A1 2003	31120	US 2002-150	353 20020517	
WO 2003099715	A1 2003	31204	WO 2003-US1	4290 20030506	
W: AE, AG,	AL, AM, AT,	AU, AZ,	BA, BB, BG, B	R, BY, BZ, CA,	CH, CN,
CO, CR,	CU, CZ, DE,	DK, DM,	DZ, EC, EE, E	S, FI, GB, GD,	GE, GH,
GM, HR,	HU, ID, IL.	IN, IS.	JP, KE, KG, K	P, KR, KZ, LC.	LK, LR,
LS, LT,	LU, LV, MA,	MD, MG,	MK, MN, MW, M	X, MZ, NO, NZ,	OM, PH,
PL, PT,	RO, RU, SC.	SD, SE,	SG, SK, SL, T	J, TM, TN. TR.	TT, TZ,
UA, UG,	US, UZ, VC,	VN, YU,	ZA, ZM, ZW, A	M, AZ, BY, KG,	KZ, MD,
RU, TJ,	TM				
RW: GH, GM,	KE, LS, MW.	MZ, SD,	SL, SZ, TZ, U	G. ZM. ZW. AT.	BE, BG,
CH, CY,	CZ, DE, DK,	EE, ES,	FI, FR, GB, G	R, HU, IE, IT,	LU, MC,
NL, PT,	RO. SE. SI.	SK, TR,	BF, BJ, CF, C	G, CI, CM, GA,	GN, GQ,
GW, ML.	MR, NE, SN,	TD, TG			
ORITY APPLN. INFO).:		US 2002-150343	A 20020517	
			UC 2002 1E02E2	A 20020E17	

PRIO

US 2002-150353 A 20020517

A method for carrying out solid state reactions under reducing conditions AR is provided. Solid state reactants include at least one inorg. metal compound and a source of reducing carbon. The reaction may be carried out in a reducing atmospheric in the presence of reducing carbon. Reducing carbon may be supplied by elemental carbon, by an organic material, or by mixts. The organic material is one that can form decomposition products containing carbon in a form capable of acting as a reductant. The reaction proceeds without significant covalent incorporation of organic material into the reaction product. In a preferred embodiment, the solid state reactants also include an alkali metal compound. The products of the method find use in lithium ion batteries as cathode active materials. Preferred active materials include lithium-transition metal phosphates and lithium-transition metal oxides. In a preferred embodiment, the reaction product contains carbon particles intimately mixed among crystals of the active materials.

Reduction IT

> (carbothermic; methods of making transition metal compds. useful as battery cathode active materials)

ΙT Phosphates, uses .

```
RL: DEV (Device component use); SPN (Synthetic preparation); PREP
    (Preparation); USES (Uses)
       (lithiated: methods of making transition metal compds. useful as
       battery cathode active materials)
    Secondary batteries
IT
        (lithjum: methods of making transition metal compds. useful as battery
       cathode active materials)
    Battery cathodes
ΙT
        (methods of making transition metal compds. useful as battery cathode
        active materials)
    Alkali metal compounds
IT
    Butadiene rubber, processes
    Carbohydrates, processes
    Hydrocarbons, processes
    RL: CPS (Chemical process); PEP (Physical, engineering or chemical
    process): PROC (Process)
        (methods of making transition metal compds. useful as battery cathode
        active materials)
    Carbonaceous materials (technological products)
    RL: CPS (Chemical process); PEP (Physical, engineering or chemical
    process); SPN (Synthetic preparation); PREP (Preparation); PROC (Process)
        (methods of making transition metal compds. useful as battery cathode
        active materials)
    Transition metal compounds
ΙT
    RL: DEV (Device component use); SPN (Synthetic preparation); PREP
     (Preparation); USES (Uses)
        (methods of making transition metal compds. useful as battery cathode
        active materials)
IT
    9003-17-2
    RL: CPS (Chemical process); PEP (Physical, engineering or chemical
    process): PROC (Process)
        (butadiene rubber, methods of making transition metal compds. useful as
        battery cathode active materials)
                                  554-13-2, Lithium carbonate
                                                                 1309-37-1.
    57-50-1, Sucrose, processes
ΙT
    Ferric oxide, processes 1313-27-5, Molybdenum oxide moo3, processes
     1333-74-0, Hydrogen, processes 1345-25-1, Ferrous oxide, processes
     7439-93-2D, Lithium, compound
    RL: CPS (Chemical process); PEP (Physical, engineering or chemical
     process); PROC (Process)
        (methods of making transition metal compds. useful as battery cathode
        active materials)
    18868-43-4P, Molybdenum dioxide 349632-76-4P, Iron lithium
    magnesium phosphate (Fe0.9LiMg0.1(PO4))
     RL: CPS (Chemical process); PEP (Physical, engineering or chemical
     process); SPN (Synthetic preparation); PREP (Preparation); PROC (Process)
        (methods of making transition metal compds. useful as battery cathode
        active materials)
     12162-92-4P. Lithium vanadium oxide liv2o5 15365-14-7P. Iron lithium
IT
     phosphate felipo4 69550-44-3P, Lithium molybdenum oxide limoo2
     84159-18-2P. Lithium vanadium phosphate Li3V2(PO4)3 108399-57-1P.
```

Lithium molybdenum oxide Li0.74MoO2 349632-79-7P, Calcium iron lithium phosphate Ca0.1Fe0.9LiPO4 507242-65-1P, Lithium molybdenum oxide Li0.85MoO2

RL: DEV (Device component use); SPN (Synthetic preparation); PREP (Preparation); USES (Uses)

(methods of making transition metal compds. useful as battery cathode active materials)

IT 11126-15-1, Lithium vanadiumoxide 37296-91-6, Lithium molybdenum oxide 39302-37-9, Lithium titanium oxide RL: MOA (Modifier or additive use): USES (Uses)

(methods of making transition metal compds. useful as battery cathode active materials) $\ \ \,$

IT 7440-44-0, Carbon, uses

RL: MOA (Modifier or additive use); USES (Uses) (particles; methods of making transition metal compds. useful as battery cathode active materials)

L2 ANSWER 6 OF 30 CAPLUS COPYRIGHT 2004 ACS on STN

ACCESSION NUMBER:

2003:717589 CAPLUS

DOCUMENT NUMBER:

139:233058

TITLE:

 \bigcirc

Alkali transition metal phosphates and related

relectrode active materials

INVENTOR(S):

Barker, Jeremy; Saidi, M. Yazid; Swoyer, Jeffrey L.

PATENT ASSIGNEE(S):

UK

SOURCE:

U.S. Pat. Appl. Publ., 14 pp.

CODEN: USXXCO

DOCUMENT TYPE:

Patent

LANGUAGE:

English

FAMILY ACC. NUM. COUNT: 1

PATENT INFORMATION:

P.A	ATENT	NO.	KIND	DATE			API	PL I C	ATIC	N NC).	DATE			
								- -			-		· -		
US	2003	170542	A1	20030	911		US	200	2-92	2317		20020	306		
WC	2003	077335	A 1	20030	918		WO	200	3-US	6998	3	20030	305		
	W:	AE, AG,	AL. AM	. AT.	AU. A	AZ. E	3A. [BB,	BG,	BR,	BY,	BZ.	CA.	CH,	CN,
		CO. CR.													
		GM. HR.		-	-										
		LS. LT.												_	
		PL. PT.			-		•								
		UA. UG.													
		RU. TJ.		, ,,,	,,,,	.0, 2	_, , ,	_,,,			, ,	υ.,	ιτα,	,	, ,
	DI.I.	GH, GM,		MiJ	м7 С	sn o	SI (S7 ·	T 7	HG	7M	714	ΔΤ	RF	RG
	IXW.														
		CH. CY.													
		NL. PT,	RO, SE	, SI,	SK, 1	TR, E	3F, 1	BJ, ∣	CF,	CG,	CI,	CM,	GA.	GN,	GQ,
		GW, ML,	MR, NE	, SN,	TD, 1	TG									
PRIORIT	Y APP	LN. INFO	. :			US	5 20	02-9	2317	,	Α .	20020	306		
AB Th	ne inv	ention c	oncerns	elect	rode	act	ive r	mate	rial	s co	ompr	ising	lit g	thium	n or
	. 1	71 7.								1.					

AB The invention concerns electrode active materials comprising lithium or other alkali metals, a transition metal, and a phosphate or similar moiety, of the formula: Aa+xMbPl-xSixO4 wherein (1) A is selected from the

group consisting of Li, Na. K, and mixts. thereof, and 0 < a < 1.0 and $0 \le x \le 1$ and (2) M comprises one or more metals, comprising at least one metal which is capable of undergoing oxidation to a higher valence state, where $0 < b \le 2$; and wherein M, a, b, and x are selected so as to maintain electroneutrality of the compound. In a preferred embodiment, M comprises at least one transition metal selected from Groups 4 to 11 of the Periodic Table. In another preferred embodiment, M comprises M'cM''d, where M' is at least one transition metal from Groups 4 to 11 of the Periodic Table; and M'' is at least one element from Groups 2, 3, 12, 13, or 14 of the Periodic Table, and c+d = b. Preferably, $0.1 \le a \le 0.8$. Preferred embodiments include those having a structure similar to the mineral olivine. This invention also provides electrodes comprising an electrode active material of this invention, and batteries that comprise a first electrode having an electrodyte.

IT Battery anodes

Battery cathodes

(alkali transition metal phosphates and related electrode active materials)

IT Oxides (inorganic), uses

RL: DEV (Device component use); USES (Uses)

(alkali transition metal phosphates and related electrode active materials)

IT Carbonaceous materials (technological products)

RL: MOA (Modifier or additive use); USES (Uses)

(alkali transition metal phosphates and related electrode active materials)

IT Chalcogenides

RL: DEV (Device component use); USES (Uses)

(metal: alkali transition metal phosphates and related electrode active materials)

IT 7440-44-0, Carbon, uses 7782-42-5, Graphite, uses

RL: DEV (Device component use); USES (Uses)

(alkali transition metal phosphates and related electrode active materials)

IT 595567-51-4P, Lithium nickel phosphate (Li0.5Ni1.25P04) 595567-52-5P 595567-53-6P, Iron lithium magnesium phosphate (FeLi0.8Mg0.1P04) 595567-55-8P 595567-59-2P, Iron lithium magnesium manganese phosphate (Fe0.5Li0.8Mg0.1Mn0.5P04) 595567-63-8P, Cobalt lithium magnesium phosphate (Co0.6Li0.5Mg0.65(P04))

RL: DEV (Device component use); SPN (Synthetic preparation); PREP (Preparation); USES (Uses)

(alkali transition metal phosphates and related electrode active materials)

L2 ANSWER 7 OF 30 CAPLUS COPYRIGHT 2004 ACS on STN

ACCESSION NUMBER:

2003:532921 CAPLUS

DOCUMENT NUMBER:

139:87848

TITLE:

Transition metal polyanion oxides as lithium ion intercalating structures for battery electrode

Print selected from Online session 22/03/2004Page 12

INVENTOR(S):

Chiang, Yet-Ming; Chung, Sung-Yoon; Bloking, Jason T.;

Andersson, Anna M.

PATENT ASSIGNEE(S):

Massachusetts Institute of Technology, USA

SOURCE:

PCT Int. Appl., 122 pp.

CODEN: PIXXD2

DOCUMENT TYPE:

Patent

LANGUAGE:

English

FAMILY ACC. NUM. COUNT: 1

PATENT INFORMATION:

```
DATE
                                          APPLICATION NO. DATE
     PATENT NO.
                     KIND
                                          _____
                                          WO 2002-US41471 20021223
     WO 2003056646
                      A1
                           20030710
        W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN,
            CO. CR. CU. CZ. DE, DK. DM. DZ. EC. EE, ES, FI, GB, GD, GE, GH,
            GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR,
            LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NO, NZ, OM, PH,
            PL, PT, RO, RU, SD, SE, SG, SK, SL, TJ, TM, TN, TR, TT, TZ, UA,
            UG, UZ, VN, YU, ZA, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM
        RW: GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZM, ZW, AT, BE, BG,
            CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL,
            PT, SE, SI, SK, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML,
            MR. NE, SN, TD, TG
                           20040108
                                          US 2002-329046
                                                           20021223
     US 2004005265
                      A1
                                       US 2001-343060P P
PRIORITY APPLN. INFO.:
                                                           20011221
                                       US 2002-388721P P
                                                           20020614
                                       US 2002-412656P P 20020920
```

The invention concerns a compound comprising a composition A.(M'1-aM'a)y(XD4)z, AB Ax.(M'1-aM'a)y(DXD4)z, or Ax.(M'1-aM'a)y(X2D7)z, and have values such that x, plus y(1-a) times a formal valence or valences of M', plus ya times a formal valence or valence of M', is equal to z times a formal valence of the XD4. X2D7, or DXD4 group; or a compound comprising a composition (A1-aM''a)xM'y(XD4)z, (A1-aM''a)xM'y(DXD4)z, (A1-aM'a)xM'y(X2D7)z and have values such that (1-a)x plus the quantity ax times the formal valence or valences of M' plus y times the formal valence or valences of M' is equal to z times the formal valence of the XD4, X2D7 or DXD4 group. In the compound. A is at least one of an alkali metal and hydrogen, M' is a first-row transition metal, X is at least one of phosphorus, sulfur, arsenic, molybdenum, and tungsten, M' any of a Group IIA, IIIA, IVA, VA, VIA, VIIA, VIIIA, IB, IIB, IIIB, IVB, VB, and VIB metal, D is at least one of oxygen, nitrogen, carbon, or a halogen, $0.0001 < a \le 0.1$, and x. y, and z are greater than zero. The compound can have a conductivity at 27° of at least about 10-8 S/cm. The compound can be a doped lithium phosphate that can intercalate lithium or hydrogen. The compound can be used in an electrochem, device including electrodes and storage batteries and can have a gravimetric capacity of at least about 80 mA-h/g while being charged/discharged at greater than about C rate of the compound

IT Battery electrodes

Electric conductivity

Fuel cell electrodes

8

```
Fuel cells
     Heat treatment
     Secondary batteries
     Solid state reaction
        (transition metal polyanion oxides as lithium ion intercalating
       structures for battery electrode)
     Intercalation compounds
IT
     RL: DEV (Device component use); USES (Uses)
       (transition metal polyanion oxides as lithium ion intercalating
        structures for battery electrode)
    7439-95-4. Magnesium, uses 7439-96-5, Manganese, uses
IT
     Niobium, uses 7440-25-7, Tantalum, uses
                                               7440-32-6, Titanium, uses
     7440-33-7, Tungsten, uses 7440-67-7, Zirconium, uses
                                                            15438-31-0, uses
     20074-52-6, uses
     RL: MOA (Modifier or additive use): USES (Uses)
        (FeLiPO4 doped with: transition metal polyanion oxides as lithium ion
        intercalating structures for battery electrode)
     554-13-2. Lithium carbonate 6047-25-2. Ferrous oxalate dihydrate
     7722-76-1. Ammonium dihydrogen phosphate
     RL: CPS (Chemical process); PEP (Physical, engineering or chemical
     process): PROC (Process)
        (transition metal polyanion oxides as lithium ion intercalating
        structures for battery electrode)
     15365-14-7P. Iron lithium phosphate felipo4
                                                  223505-09-7P. Iron lithium
ΙT
                                       478819-82-8P, Iron lithium titanium
     titanium phosphate 411234-54-3P
     phosphate FeLi0.99Ti0.01(PO4) 478819-83-9P, Iron lithium niobium
     phosphate FeLi0.99Nb0.01(PO4) 478819-84-0P, Iron lithium
     magnesium phosphate FeLio 99MgO 01(PO4) 478819-85-1P, Aluminum iron
    Tithium phosphate (Alo.01FeLio.99(PO4)) 478819-87-3P, Iron lithium
     titanium phosphate Fe0.99LiTi0.01(PO4) 478819-89-5P, Iron lithium
                                                496816-58-1P, Iron lithium
     zirconium phosphate (Fe0.99LiZr0.01(P04))
     zirconium phosphate (Fe0.98LiZr0.02(PO4))
                                                531493-25-1P. Iron lithium
     titanium phosphate Fe0.98LiTi0.02(P04)
                                            554453-36-0P 554453-37-1P
     554453-38-2P
                   554453-39-3P
                                  554453-40-6P
                                                 554453-41-7P
                   554453-44-0P. Iron lithium zirconium phosphate
     554453-42-8P
                           554453-45-1P. Iron lithium niobium phosphate
     (Fe0.95LiZr0.05(P04))
     (FeLi0.98Nb0.02(PO4))
                            554453-46-2P, Iron lithium niobium phosphate
                            554453-47-3P, Iron lithium phosphate
     (FeLi0.96Nb0.04(PO4))
                          554453-48-4P. Iron lithium phosphate
     (Fe1.01Li0.99(PO4))
     (Fe0.99Li1.01(PO4))
     RL: DEV (Device component use); PRP (Properties); SPN (Synthetic
     preparation); PREP (Preparation); USES (Uses)
        (transition metal polyanion oxides as lithium ion intercalating
        structures for battery electrode)
    7440-37-1, Argon, uses 7727-37-9, Nitrogen, uses
IT
     RL: TEM (Technical or engineered material use); USES (Uses)
        (transition metal polyanion oxides as lithium ion intercalating
        structures for battery electrode)
                              THERE ARE 8 CITED REFERENCES AVAILABLE FOR THIS
REFERENCE COUNT:
```

RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

DEL 2 ANSWER 8 OF 30 CAPLUS COPYRIGHT 2004 ACS on STN

ACCESSION NUMBER:

2003:125703 CAPLUS

DOCUMENT NUMBER:

138:388269

TITLE:

HT-XRD study of synthetic ferrian magnesian spodumene:

the effect of site dimension on the P21/c→C2/c

phase transition

AUTHOR(S):

Camara, F.; Iezzi, G.; Oberti, R.

CORPORATE SOURCE:

Istituto di Geoscienze e Georisorse, CNR, Pavia,

27100. Italy

SOURCE:

Physics and Chemistry of Minerals (2003), 30(1), 20-30

CODEN: PCMIDU; ISSN: 0342-1791

PUBLISHER:

Springer-Verlag

DOCUMENT TYPE:

Journal

LANGUAGE:

English

Ferrian magnesian spodumene was synthesized in the MgO-Li2O-FeO-SiO2-H2O system at 0.4 GPa, 700° , with f02 = NNO + 2.3. The space group at room temperature is P21/c [a = 9.638(3), b = 8.709(2), c = 5.258(2) Å. $\beta = 109.83(3)$. volume 415.2 Å3]. The structure is topol. equivalent to that of ferrian spodumene, LiFeSi206, and has two sym. independent tetrahedral chains, A and B, and two independent octahedral sites, M1 and M2. The crystal-chemical composition was determined combining electron microprobe. SIMS and single-crystal XRD anal., yielding M2(Li0.85Mg0.09Fe2+0.06) M1(Fe3+0.85Mg0.15)Si206. Li is ordered at the M2 site and Fe3+ is ordered at the M1 site, whereas Mg (and Fe2+) distribute over both octahedral sites. Structure refinements done at different temps. (25, 70, 95, 125. 150 and 200 °C) allowed characterization of a reversible displacive P21/c→ C2/c transition at 106°. Previous HT-XRD studies of Li-clinopyroxenes had shown that the transition temperature is inversely related to the size of the M1 cation. For the crystal of this work, the aggregate ionic radius at M1 is longer than that of ferrian spodumene, for which the transition temperature is -44°. The higher transition temperature observed can only be explained on the basis of the shorter aggregate radius at the M2 site (due to the presence of Mg substituting after Li). in keeping with the results obtained for ferromagnesian P21/c pyroxenes. The effects of all the chemical substitutions must be considered when modeling transition temps, and thermodn, behavior in clinopyroxenes.

IT Structural phase transition

(high-temperature XRD study of synthetic magnesian iron spodumene showing effect of site dimension on the $P21/c \rightarrow C2/c$ phase transition)

IT Bond angle

Bond length

(in synthetic magnesian iron spodumene)

IT Order

(in synthetic magnesian iron spodumene undergoing P21/c \rightarrow C2/c phase transition at 106°)

IT Crystal structure

(of synthetic magnesian iron spodumene undergoing P21/c \rightarrow C2/c phase transition at 106°)

IT 528603-23-8, Iron lithium magnesium silicate

(Fe0.91Li0.85Mg0.24(Si03)2)

RL: PRP (Properties)

(high-temperature XRD study of synthetic magnesian iron spodumene showing effect of site dimension on the P21/c→C2/c phase transition)

REFERENCE COUNT:

30

THERE ARE 30 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

L2 ANSWER 9 OF 30 CAPLUS COPYRIGHT 2004 ACS on STN

ACCESSION NUMBER:

2003:83100 CAPLUS

DOCUMENT NUMBER:

138:341011

TITLE:

Lithium iron(II) phospho-olivines prepared by a novel

carbothermal reduction method

AUTHOR(S):

Barker, J.; Saidi, M. Y.; Swoyer, J. L.

CORPORATE SOURCE:

Valence Technology Incorporated, Henderson, NV, 89015.

USA

SOURCE:

Electrochemical and Solid-State Letters (2003), 6(3),

A53-A55

CODEN: ESLEF6; ISSN: 1099-0062

PUBLISHER:

Electrochemical Society

DOCUMENT TYPE: LANGUAGE: Journal English

AB The electroactive materials LiFePO4 and LiFe0.9Mg0.1PO4 have been synthesized by a novel carbothermal reduction (CTR) method. The transition metal reduction and lithium incorporation processes are each facilitated by the high temperature carbothermal reaction based on the C→CO transition. These CTR conditions favor stabilization of the iron as Fe2+ as well as offering some control of the product morphol. and conductivity Electrochem. evaluation of the CTR LiFe0.9Mg0.1PO4 reveals a lithium insertion plateau around 3.4 V vs. Li together with a specific capacity of over 150 mAh/g. Differential capacity data confirm the two-phase nature of the insertion reactions as well as the outstanding ionic reversibility. Few tech. obstacles have been encountered in scaling the CTR process from a laboratory

process to that required for pilot production

IT Reduction

IT Battery electrodes

IT Secondary batteries

(lithium; lithium iron(II) phospho-olivines prepared by novel carbothermal reduction method)

IT 15365-14-7P, Iron lithium phosphate FeLiPO4 **349632-76-4P**, Iron lithium magnesium phosphate (Fe0.9LiMg0.1(PO4))

RL: PRP (Properties); SPN (Synthetic preparation); TEM (Technical or engineered material use); PREP (Preparation); USES (Uses)

(battery electrodes; lithium iron(II) phospho-olivines prepared by novel carbothermal reduction method)

REFERENCE COUNT:

THERE ARE 19 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

ANSWER 10 OF 30 CAPLUS COPYRIGHT 2004 ACS on STN L2

ACCESSION NUMBER:

2002:928099 CAPLUS

DOCUMENT NUMBER:

138:6481

TITLE:

Process for producing carbon-containing lithium-iron composite phosphorus oxide for lithium secondary

battery cathode active material .

INVENTOR(S):

Kohzaki, Masao; Takeuchi, Youji; Ukyo, Yoshio Kabushiki Kaisha Toyota Chuo Kenkyusho, Japan

U.S. Pat. Appl. Publ., 11 pp.

SOURCE:

CODEN: USXXCO

DOCUMENT TYPE:

Patent

LANGUAGE:

English

FAMILY ACC. NUM. COUNT: 1

PATENT ASSIGNEE(S):

PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
US 2002182497	A1	20021205	US 2002-143946	20020514
JP 2003034534	A2	20030207	JP 2002-132603	20020508
PRIORITY APPLA INFO			JP 2001-145396 A	20010515

AB A carbon-containing lithium-iron composite phosphorus oxide for a lithium secondary battery pos. electrode active material, includes particles being composed of a lithium-iron composite phosphorus oxide having an olivine structure whose basic composition is LiFePO4, and being composited with carbonaceous fine particles. A process for producing the same includes the steps of mixing a lithium compound making a lithium source, an iron compound making an iron source, a phosphorus-containing ammonium salt making a phosphorus source and carbonaceous fine particles, thereby preparing a mixture, and calcicing the mixture at a temperature of from 600° or more to 750° or less.

Carbon black, uses

RL: MOA (Modifier or additive use); USES (Uses)

(composited with lithium iron phosphate; process for producing carbon-containing lithium-iron composite phosphorus oxide for lithium secondary battery cathode active material)

IT Secondary batteries

> (lithium; process for producing carbon-containing lithium-iron composite phosphorus oxide for lithium secondary battery cathode active material)

IT Battery cathodes

> (process for producing carbon-containing lithium-iron composite phosphorus oxide for lithium secondary battery cathode active material)

15365-14-7P, Iron lithium phosphate FeLiPO4 476669-99-5P, Iron lithium ΙT manganese phosphate (Fe0.8-0.98LiMn0.02-0.2(P04)) 476670-01-6P.

Iron lithium magnesium phosphate (Fe0.8-0.98LiMg0.02-0.2(PO4))

476670-03-8P. Iron lithium nickel phosphate (Fe0.8-0.98LiNi0.02-0.2(PO4))

476670-05-0P, Cobalt iron lithium phosphate (Co0.02-0.2Fe0.8-0.98Li(PO4))

476670-07-2P, Copper iron lithium phosphate (Cu0.02-0.2Fe0.8-0.98Li(PO4))

476670-10-7P. Iron lithium zinc phosphate (Fe0.8-0.98LiZn0.02-0.2(P04))

476670-12-9P, Germanium iron lithium phosphate (GeO.02-0.2FeO.8-

0.98Li(PO4))

Battery cathodes

0

```
RL: DEV (Device component use); SPN (Synthetic preparation); PREP
    (Preparation): USES (Uses)
       (carbon composited; process for producing carbon-containing lithium-iron
       composite phosphorus oxide for lithium secondary battery cathode active
       material)
    7440-44-0. Carbon, uses
    RL: MOA (Modifier or additive use); USES (Uses)
       (composited with lithium iron phosphate; process for producing
       carbon-containing lithium-iron composite phosphorus oxide for lithium
       secondary battery cathode active material)
    516-03-0. Ferrous oxalate
                                           7722-76-1. Ammonium dihydrogen
                                554-13-2
     phosphate
    RL: CPS (Chemical process); PEP (Physical, engineering or chemical
    process); PROC (Process)
       (process for producing carbon-containing lithium-iron composite phosphorus
       oxide for lithium secondary battery cathode active material)
    412351-36-1P, Iron lithium manganese phosphate (Fe0.9LiMn0.1(PO4))
     476670-14-1P, Iron lithium manganese phosphate (Fe0.85LiMn0.15(PO4))
     RL: DEV (Device component use); SPN (Synthetic preparation); PREP
     (Preparation); USES (Uses)
        (process for producing carbon-containing lithium-iron composite phosphorus
       oxide for lithium secondary battery cathode active material)
L2 ANSWER 11 OF 30 CAPLUS COPYRIGHT 2004 ACS on STN
ACCESSION NUMBER:
                         2002:794447 CAPLUS
                         138:41938
DOCUMENT NUMBER:
                        Electronically conductive phospho-olivines as lithium
TITLE:
                         storage electrodes
                         Chung, Sung-Yoon; Bloking, Jason T.; Chiang, Yet-Ming
AUTHOR(S):
                         Department of Materials Science and Engineering.
CORPORATE SOURCE:
                        Massachusetts Institute of Technology, Cambridge, MA,
                         02139. USA
                         Nature Materials (2002), 1(2), 123-128
SOURCE:
                         CODEN: NMAACR; ISSN: 1476-1122
                         Nature Publishing Group
PUBLISHER:
DOCUMENT TYPE:
                         Journal
                       English
LANGUAGE:
    Lithium transition metal phosphates are of interest as storage cathodes
     for rechargeable Li batteries because of their high energy d., low raw
     materials cost, environmental friendliness and safety. Their key
     limitation was extremely low electronic conductivity, believed to be intrinsic to
     this family of compds. Controlled cation nonstoichiometry combined with
     solid-solution doping by metals supervalent to Li+ increases the electronic
     conductivity of LiFePO4 by a factor of .apprx.108. The resulting materials show
     near-theor. energy d. at low charge/discharge rates, and retain
     significant capacity with little polarization at rates as high as 6,000
     mA/g. In a conventional cell design, they may allow development of Li
     batteries with the highest power d. yet.
```

```
(electronically conductive phospho-olivines as lithium storage cathodes
        for batteries)
     15365-14-7. Iron lithium phosphate (FeLiPO4)
IT
     RL: DEV (Device component use); PRP (Properties); USES (Uses)
        (doped: electronically conductive phospho-olivines as lithium storage
        cathodes for batteries)
     478819-81-7. Iron lithium zirconium phosphate (FeLi0.99Zr0.01(PO4))
IT
     478819-82-8. Iron lithium titanium phosphate (FeLi0.99Ti0.01(PO4))
     478819-83-9. Iron lithium niobium phosphate (FeLi0.99Nb0.01(PO4))
     478819-84-0, Iron lithium magnesium phosphate
                            478819-85-1. Aluminum iron lithium phosphate
     (FeLi0.99Mg0.01(PO4))
                            478819-86-2. Iron lithium niobium phosphate
     (Al0.01FeLi0.99(PO4))
                            478819-87-3, Iron lithium titanium phosphate
     (Fe0.99LiNb0.01(PO4))
                            478819-89-5, Iron lithium zirconium phosphate
     (Fe0.99LiTi0.01(PO4))
                            478819-90-8. Aluminum iron lithium phosphate
     (Fe0.99LiZr0.01(PO4))
     (AlO.01Fe0.99Li(PO4)) 478819-92-0, Iron lithium magnesium
     phosphate (Fe0.99LiMq0.01(P04))
     RL: DEV (Device component use); PRP (Properties); USES (Uses)
        (electronically conductive phospho-olivines as lithium storage cathodes
        for batteries)
                         29
                               THERE ARE 29 CITED REFERENCES AVAILABLE FOR THIS
REFERENCE COUNT:
                               RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT
    ANSWER 12 OF 30 CAPLUS COPYRIGHT 2004 ACS on STN
                         2002:573514 CAPLUS
ACCESSION NUMBER:
                         137:143023
DOCUMENT NUMBER:
                         secondary nonaqueous electrolyte battery
TITLE:
                         Nakamura, Masaya: Saito, Hirohiko
INVENTOR(S):
                         Denso Co., Ltd., Japan
PATENT ASSIGNEE(S):
SOURCE:
                         Jpn. Kokai Tokkyo Koho, 7 pp.
                         CODEN: JKXXAF
DOCUMENT TYPE:
                         Patent
LANGUAGE:
                         Japanese
FAMILY ACC. NUM. COUNT:
PATENT INFORMATION:
     PATENT NO.
                      KIND
                           DATE
                                           APPLICATION NO.
                       A2
                            20020802
                                           JP 2001-13581
                                                            20010122
     JP 2002216770
PRIORITY APPLN. INFO.:
                                        JP 2001-13581
                                                            20010122
    The battery is a Li battery using an olivine structured LiMxFel-xPO4 [ M =
     metal(s) excluding Fe; x = 0-0.5] cathode active mass and contains a
     conductive control material, which cuts off elec. current in the battery
     when the conductivity of the cathode and/or anode drops sharply due to
     electrochem. doping and undoping. The conductive control material is
```

preferably a material that can be p-doped, e.g., polyaniline or

polypyrrole, and is in the cathode. The battery using the conductive control material and the cathode active mass has a high safety when

IT Battery cathodes

overcharging and in hot environment.

(cathodes containing lithium iron phosphate in secondary lithium batteries containing conductivity control materials)

Safety IT

(conductivity control materials in secondary lithium battery using cathode containing lithium iron phospahte for safety)

Polyanilines IT

RL: DEV (Device component use); USES (Uses)

(conductivity control materials in secondary lithium battery using cathode containing lithium iron phospahte for safety)

15365-14-7. Iron lithium phosphate (FeLiPO4) 113066-78-7. Cobalt lithium nickel oxide (Co0.4LiNi0.602) 155472-68-7, Lithium manganese oxide

(Li1.1Mn1.904) 190436-16-9 **349632-85-5** 444816-41-5

444816-42-6. Aluminum iron lithium phosphate (Al0.2Fe0.8Li(PO4))

444816-45-9 444816-47-1 444816-43-7 444816-44-8

RL: DEV (Device component use); USES (Uses)

(cathodes containing lithium iron phosphate in secondary lithium batteries containing conductivity control materials)

25233-30-1. Polyaniline

RL: DEV (Device component use); TEM (Technical or engineered material

use): USES (Uses)

(conductivity control materials in secondary lithium battery using cathode containing lithium iron phospahte for safety)

L2 ANSWER 13 OF 30 CAPLUS COPYRIGHT 2004 ACS on STN

ACCESSION NUMBER:

2002:428819 CAPLUS

DOCUMENT NUMBER:

137:8642

TITLE:

Methods of making lithium metal compounds useful as

cathode active materials in batteries

INVENTOR(S):

Barker, Jeremy; Yazid, Saidi M.; Swoyer, Jeffrey L. Valence Technology, Inc., USA

PATENT ASSIGNEE(S):

PCT Int. Appl., 85 pp.

SOURCE:

CODEN: PIXXD2

DOCUMENT TYPE:

Patent

LANGUAGE:

English

FAMILY ACC. NUM. COUNT: 1

PATENT INFORMATION:

PATENT NO.	KIND DATE	APPLICATION NO. DATE
WO 2002044084	A2 20020606	WO 2001-US43633 20011119
W: AE, AG,	AL, AM, AT, AU,	AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN,
CO, CR,	CU, CZ, DE, DK,	DM. DZ. EC. EE. ES. FI. GB. GD. GE. GH.
GM, HR,	HU, ID, IL, IN,	IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR,
LS, LT,	LU, LV, MA, MD,	MG, MK, MN, MW, MX, MZ, NO, NZ, OM, PH,
PL, PT,	RO, RU, SD, SE,	SG, SI, SK, SL, TJ, TM, TR, TT, TZ, UA,
UG, US,	UZ, VN, YU, ZA,	ZM, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM
RW: GH, GM,	KE, LS, MW, MZ,	SD, SL, SZ, TZ, UG, ZM, ZW, AT, BE, CH,
CY, DE,	DK, ES, FI, FR,	GB, GR, IE, IT, LU, MC, NL, PT, SE, TR,
BF, BJ,	CF, CG, CI, CM,	GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG
US 6645452	B1 20031111	US 2000-724085 20001128

AU 2002-17799 Α5 20020611 20011119 AU 2002017799 EP 2001-998506 20011119 A2 20030917 EP 1343720 R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT, IE. SI. LT. LV. FI, RO. MK, CY, AL, TR PRIORITY APPLN. INFO.: US 2000-724085 A1 20001128 WO 2001-US43633 W 20011119

AB The invention provides a novel method for making lithium mixed metal materials for battery cathodes. The lithium mixed metal materials comprise lithium and at least one other metal besides lithium. The invention involves the reaction of a metal compound, a phosphate compound, with a reducing agent to reduce the metal and form a metal phosphate. The invention also includes methods of making lithium metal oxides involving reaction of a lithium compound and a metal oxide with a reducing agent.

IT Reduction

(carbothermic; methods of making lithium metal compds. useful as cathode active materials in batteries)

IT Secondary batteries

(lithium; methods of making lithium metal compds. useful as cathode active materials in batteries)

IT Battery cathodes

Thermite process

(methods of making lithium metal compds. useful as cathode active materials in batteries)

TT 7664-38-2D. Phosphoric acid. transition metal compds. 7722-76-1.

Ammonium dihydrogen phosphate 7757-87-1. Magnesium phosphate mg3(po4)2
7779-90-0. Zinc phosphate zn3(po4)2 7783-28-0. Diammonium hydrogen
phosphate 7789-04-0. Chromium phosphate crpo4 7789-24-4. Lithium
fluoride, processes 10045-86-0. Iron phosphate fepo4 13453-80-0.

Lithium dihydrogen phosphate 14154-09-7. Manganese phosphate Mn3(PO4)2
14940-41-1. Iron phosphate fe3(po4)2 70172-55-3. Titanium phosphate
tipo4

RL: CPS (Chemical process); PEP (Physical, engineering or chemical process); PROC (Process)

(methods of making lithium metal compds. useful as cathode active materials in batteries)

7664-38-2DP. Phosphoric acid, lithiated transition metal compds. 12162-92-4P, Lithium vanadium oxide liv2o5 15365-14-7P, Iron lithium phosphate felipo4 84159-18-2P, Lithium vanadium phosphate Li3V2(PO4)3 349632-76-4P. Iron lithium magnesium phosphate (Fe0.9LiMg0.1(PO4)) 372075-82-6P. Lithium manganese fluoride phosphate LiMnFPO4 372075-83-7P. Lithium vanadium fluoride phosphate (LiVF(PO4)) 372075-84-8P. Chromium lithium fluoride phosphate CrLiFPO4 372075-85-9P. Lithium titanium fluoride phosphate LiTiFPO4 372075-86-0P 372075-87-1P, Iron lithium fluoride phosphate FeLiFPO4 433708-98-6P. Copper lithium fluoride phosphate (CuLiF(PO4)) 433708-99-7P, Cobalt 433709-00-3P. Lithium nickel lithium fluoride phosphate (CoLiF(PO4)) fluoride phosphate (LiNiF(PO4)) 433709-01-4P RL: DEV (Device component use); SPN (Synthetic preparation); PREP

RL: DEV (Device component use); SPN (Synthetic preparation); PREP (Preparation); USES (Uses)

(methods of making lithium metal compds. useful as cathode active

materials in batteries)

IT 1333-74-0, Hydrogen, reactions

RL: RCT (Reactant); RACT (Reactant or reagent)

(methods of making lithium metal compds. useful as cathode active materials in batteries)

IT 124-38-9, Carbon dioxide, uses 630-08-0, Carbon monoxide, uses

7440-37-1. Argon. uses 7727-37-9. Nitrogen, uses

RL: TEM (Technical or engineered material use); USES (Uses) (methods of making lithium metal compds. useful as cathode active

materials in batteries)

IT 7440-44-0, Carbon, reactions

RL: RCT (Reactant); RACT (Reactant or reagent)

(reducing agent; methods of making lithium metal compds. useful as cathode active materials in batteries)

L2 ANSWER 14 OF 30 CAPLUS COPYRIGHT 2004 ACS on STN

ACCESSION NUMBER:

2002:292205 CAPLUS

DOCUMENT NUMBER:

136:328153

TITLE:

Nonaqueous electrolyte battery containing complex

lithium oxide positive electrode with olivine crystal

structure

INVENTOR(S):

Fukushima, Gen; Hosoya, Mamoru; Hisayama, Junji; Takahashi, Kimio; Sato, Atsushi; Okawa, Takeshi

PATENT ASSIGNEE(S):

Sony Corp., Japan

SOURCE:

Jpn. Kokai Tokkyo Koho, 9 pp.

CODEN: JKXXAF

DOCUMENT TYPE:

Patent

LANGUAGE:

Japanese

FAMILY ACC. NUM. COUNT: 1

PATENT INFORMATION:

PATENT NO. KIND DATE APPLICATION NO. DATE

JP 2002117903 A2 20020419 JP 2000-306878 20001005
PRIORITY APPLN. INFO.: JP 2000-306878 20001005

AB A battery comprises a neg. electrode, a pos. electrode from a compound LixFel-yMyPO4 (M is Mn, Cr, Co, Cu, Ni, V, Mo, Ti, Zn, Al, Ga, Mg, B, Nb; x = 0.05-1.2, yr = 0-0.8), and a separator. The pos. electrode compound has an olivine crystal structure. The combined volume (a) of the electrodes and volume (b) of separator satisfy the following conditions: $0.17 \le b/(a + b) \le 0.39$. The battery has improved balance of energy d.

IT Secondary battery separators

(controlled volume of; nonaq. electrolyte battery containing complex lithium oxide pos. electrode with olivine crystal structure)

IT Secondary batteries

(lithium; nonaq. electrolyte battery containing complex lithium oxide pos. electrode with olivine crystal structure)

IT Battery electrodes

(nonaq. electrolyte battery containing complex lithium oxide pos. electrode with olivine crystal structure)

Olivine-group minerals

RL: DEV (Device component use); USES (Uses)

```
Olivine-group minerals
    RL: DEV (Device component use); USES (Uses)
       (nonag. electrolyte battery containing complex lithium oxide pos. electrode
       with olivine crystal structure)
ΙT
    Crystal structure
       (olivine; nonaq. electrolyte battery containing complex lithium oxide pos.
       electrode with olivine crystal structure)
ΙT
    15365-14-7
                 407629-83-8
                              407629-87-2 407629-90-7
                                                           407629-95-2
    407630-01-7
                  407630-05-1
                                407630-10-8 407630-14-2
                                                            407630-25-5.
    Aluminum iron lithium phosphate (Al0.7Fe0.3Li(PO4))
                                                          407630-29-9, Gallium
    iron lithium phosphate (GaO.7FeO.3Li(PO4)) 407630-35-7
    407630-40-4, Boron iron lithium phosphate (B0.75Fe0.25Li(PO4))
    407630-46-0 412267-84-6, Iron lithium zinc phosphate (Fe0.3LiZn0.7(P04))
    RL: DEV (Device component use); USES (Uses)
       (pos. electrode containing; nonag, electrolyte battery containing complex
       lithium oxide pos. electrode with olivine crystal structure)
    ANSWER 15 OF 30 CAPLUS COPYRIGHT 2004 ACS on STN
ACCESSION NUMBER:
                        2002:292166 CAPLUS
DOCUMENT NUMBER:
                        136:328120
                        Nonaqueous electrolyte secondary battery with lithium
TITLE:
                        complex oxide positive electrode mix additionally
                        containing lithium carbonate for improved
                        charge-discharge cycle characteristic at high
                        temperature
                        Hosoya, Mamoru; Fukushima, Gen
INVENTOR(S):
                        Sony Corp., Japan
PATENT ASSIGNEE(S):
                         Jpn. Kokai Tokkyo Koho, 14 pp.
SOURCE:
                        CODEN: JKXXAF
DOCUMENT TYPE:
                        Patent
LANGUAGE:
                        Japanese
FAMILY ACC. NUM. COUNT:
PATENT INFORMATION:
                                          APPLICATION NO.
    PATENT NO.
                     KIND DATE
                                          -----
    JP 2002117843
                      A2
                           20020419
                                          JP 2000-306875
                                                           20001005
PRIORITY APPLN. INFO.:
                                       JP 2000-306875
                                                           20001005
    A nonag. electrolyte secondary battery comprises a LixFe1-yMyP04 (M is Mn.
    Cr. Co. Cu. Ni. V. Mo. Ti. Zn. Al. Ga. Mg. B. and/or Nb; x = 0.05-1.2. yr
    ≤0.8), pos. electrode active mix addnl. containing Li2CO3. The battery
    has improved charge-discharge cycle characteristic at high temperature
    Battery cathodes
IT
       (nonag. electrolyte secondary battery with lithium complex oxide pos.
       electrode mix addnl. containing lithium carbonate for improved
       charge-discharge cycle characteristic at high temperature)
```

(nonag. electrolyte secondary battery with lithium complex oxide pos.

electrode mix addnl. containing lithium carbonate for improved

```
charge-discharge cycle characteristic at high temperature)
     15365-14-7. Iron lithium phosphate (FeLiPO4) 407629-83-8 407629-87-2
ΙT
                                407630-01-7 407630-05-1
                                                            407630-10-8
     407629-90-7
                  407629-95-2
                  407630-25-5, Aluminum iron lithium phosphate
     407630-14-2
     (AlO.7Fe0.3Li(PO4)) 407630-29-9, Gallium iron lithium phosphate
     (Ga0.7Fe0.3Li(PO4)) 407630-35-7 407630-40-4, Boron iron
     lithium phosphate (B0.75Fe0.25Li(PO4)) 407630-46-0
                                                           412267-84-6. Iron
     lithium zinc phosphate (Fe0.3LiZn0.7(PO4))
     RL: DEV (Device component use); USES (Uses)
        (nonag. electrolyte secondary battery with lithium complex oxide pos.
       electrode mix addnl. containing lithium carbonate for improved
        charge-discharge cycle characteristic at high temperature)
     554-13-2. Lithium carbonate
     RL: DEV (Device component use); MOA (Modifier or additive use); USES
     (Uses)
        (nonag. electrolyte secondary battery with lithium complex oxide pos.
        electrode mix addnl. containing lithium carbonate for improved
       charge-discharge cycle characteristic at high temperature)
    ANSWER 16 OF 30 CAPLUS COPYRIGHT 2004 ACS on STN
                        2002:272915 CAPLUS
ACCESSION NUMBER:
DOCUMENT NUMBER:
                        136:297401
                        Nonaqueous electrolyte battery with high discharge
TITLE:
                        capacity
                         Sakai, Hidecki; Fukushima, Yuzuru; Kuyama, Junji;
INVENTOR(S):
                        Hosoya, Mamoru
PATENT ASSIGNEE(S):
                         Sony Corporation, Japan
                         Eur. Pat. Appl., 17 pp.
SOURCE:
                        CODEN: EPXXDW
DOCUMENT TYPE:
                        Patent
LANGUAGE:
                        English
FAMILY ACC. NUM. COUNT:
PATENT INFORMATION:
     PATENT NO.
                                          APPLICATION NO.
                                                           DATE
                      KIND DATE
                            ----------
                                          _____
     ------
                      ----
     EP 1195838
                      A2
                           20020410
                                          EP 2001-123895
                                                           20011005
         R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT,
             IE, SI, LT, LV, FI, RO
     JP 2002117908
                      A2
                           20020419
                                          JP 2000-308303
                                                           20001006
     CN 1348230
                      Α
                            20020508
                                          CN 2001-130350
                                                           20010930
                                          US 2001-971912
     US 2002150816 ·
                      A1
                            20021017
                                                           20011005
     TW 523945
                       В
                            20030311
                                          TW 2001-90124693 20011005
                                        JP 2000-308303
                                                       A 20001006
PRIORITY APPLN. INFO.:
     A nonag. electrolyte cell is disclosed having high discharge capacity, an
     improved capacity upkeep ratio and optimum cyclic characteristics. The
     nonaq. electrolyte cell has a cell device including a strip-shaped cathode
```

material and a strip-shaped anode material, layered and together via a

separator and coiled a plural number of times, a nonaq. electrolyte solution, and a cell can for accommodating cell device and the nonaq. electrolyte

```
solution The cathode employs a cathode active material containing a compound of
    the olivinic structure represented by the general formula LixFel-yMyPO4.
    where M is at least one selected from the group consisting of Mn. Cr. Co.
    Cu. Ni, V. Mo, Ti. Zn. Al. Ga. Mg. B and Nb. with 0.05 \le x \le x \le 10^{-5}
    1.2 and 0 \le y \le 0.8, with the compound being used either
    singly or in combination with other materials. The ratio of an inner
    diameter d to an outer diameter D of cell device is selected so that 0.05 < d/D
    < 0.5.
    Secondary batteries
IT
        (lithium; nonaq. electrolyte battery with high discharge capacity)
IT
    Battery cathodes
        (nonag. electrolyte battery with high discharge capacity)
ΙT
    Carbon black, uses
    RL: DEV (Device component use); USES (Uses)
        (nonag. electrolyte battery with high discharge capacity)
    Lithium alloy, base
·ΙΤ
    RL: DEV (Device component use); USES (Uses)
        (nonag. electrolyte battery with high discharge capacity)
    7439-93-2, Lithium, uses 15365-14-7, Iron lithium phosphate felipo4
IT
    407606-22-8, Chromium iron lithium phosphate (Cr0-0.8Fe0.2-1Li0.05-
                407606-24-0, Cobalt iron lithium phosphate
     (CoO-0.8Fe0.2-1Li0.05-1.2(PO4))
                                       407606-26-2, Copper iron lithium
                                               407606-28-4, Aluminum iron
     phosphate (Cu0-0.8Fe0.2-1Li0.05-1.2(PO4))
     lithium phosphate (AlO-0.8Fe0.2-1Li0.05-1.2(PO4))
                                                         407606-30-8. Gallium
     iron lithium phosphate (GaO-0.8Fe0.2-1Li0.05-1.2(PO4))
                                                              407606-32-0.
     Boron iron lithium phosphate (B0-0.8Fe0.2-1Li0.05-1.2(PO4))
                                                                  407606-34-2.
     Iron lithium manganese phosphate (Fe0.2-1Li0.05-1.2Mn0-0.8(P04))
     407606-36-4, Iron lithium nickel phosphate (Fe0.2-1Li0.05-1.2Ni0-0.8(PO4))
     407606-39-7, Iron lithium vanadium phosphate (Fe0.2-1Li0.05-1.2V0-
     0.8(P04)
                407606-42-2. Iron lithium molybdenum phosphate
     (Fe0.2-1Li0.05-1.2Mo0-0.8(PO4))
                                      407606-44-4. Iron lithium titanium
     phosphate (Fe0.2-1Li0.05-1.2Ti0-0.8(PO4)) 407606-47-7, Iron lithium zinc
     phosphate (Fe0.2-1Li0.05-1.2Zn0-0.8(P04)) 407606-49-9, Iron
     lithium magnesium phosphate (Fe0.2-1Li0.05-1.2Mg0-0.8(PO4))
                                                                   407606-51-3.
     Iron lithium niobium phosphate (Fe0.2-1Li0.05-1.2Nb0-0.8(PO4))
     407629-83-8
                  407629-87-2
                                407629-90-7
                                               407629-95-2
                                                             407630-01-7
     407630-05-1
                  407630-10-8
                                 407630-14-2
                                               407630-25-5. Aluminum iron
     lithium phosphate (Alo.7Fe0.3Li(PO4)) 407630-29-9, Gallium iron lithium
     phosphate (Ga0.7Fe0.3Li(PO4)) 407630-35-7
                                                 407630-40-4, Boron
     iron lithium phosphate (B0.75Fe0.25Li(PO4))
                                                   408501-54-2
     RL: DEV (Device component use); USES (Uses)
        (nonag. electrolyte battery with high discharge capacity)
    ANSWER 17 OF 30 CAPLUS COPYRIGHT 2004 ACS on STN
ACCESSION NUMBER:
                         2002:272914 CAPLUS
DOCUMENT NUMBER:
                         136:297400
                         Nonaqueous electrolyte secondary battery using
TITLE:
                         olivinic lithium phosphorus oxide cathode active
```

Okawa, Tsuyoshi; Hosoya, Mamoru; Kuyama, Junji;

INVENTOR(S):

Fukushima, Yuzuru

PATENT ASSIGNEE(S):

Sony Corporation, Japan Eur. Pat. Appl., 15 pp.

SOURCE:

CODEN: EPXXDW

DOCUMENT TYPE:

Patent English

LANGUAGE:

. 1

FAMILY ACC. NUM. COUNT: 1

PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
EP 1195837	A2	20020410	EP 2001-123893	20011005
R: AT, BE,	CH, DE	, DK, ES,	FR, GB, GR, IT, LI, LU	, NL, SE, MC, PT,
IE, SI,	LT, LV	, FI, RO		
JP 2002117907	A2	20020419	JP 2000-308302	20001006
CN 1350342	Α	20020522	CN 2001-139303	20010930
US 2002106563	A1	20020808	US 2001-972375	20011005
PRIORITY APPLN. INFO	. :		JP 2000-308302 A	20001006

AB In a battery, liquid leakage or destruction may be prevented as the apparent energy d. per unit volume of the cell is maintained. The cell uses, as a cathode active material, a compound of an olivinic crystal structure having the formula LixFel-xMyPO4, where M is at least one selected from the group of Mn, Cr, Co, Cu, Ni, V, Mo, Ti, Zn, Al, Ga, Mg, B and Nb and 0.05 $\leq x \leq 1.2$ and $0 \leq y \leq 0.8$. By adjusting the amount of the electrolyte solution, the amount of the void in the container is set so as to be not less than 0.14 mL and not more than 3.3 mL per 1 Ah of the cell capacity.

`IT Secondary batteries

(lithium; nonaq. electrolyte secondary battery using olivinic lithium phosphorus oxide cathode active material)

IT Battery cathodes

Composites

(nonaq. electrolyte secondary battery using olivinic lithium phosphorus oxide cathode active material)

IT Coke

RL: DEV (Device component use): USES (Uses)

(pitch; nonaq. electrolyte secondary battery using olivinic lithium phosphorus oxide cathode active material)

IT 108-32-7, Propylene carbonate 616-38-6. Dimethyl carbonate 7440-44-0. Carbon, uses 15365-14-7, Iron lithium phosphate felipo4 21324-40-3, Lithium hexafluorophosphate 407606-22-8. Chromium iron lithium phosphate (Cr0-0.8Fe0.2-1Li0.05-1.2(PO4)) 407606-24-0. Cobalt iron lithium phosphate (Co0-0.8Fe0.2-1Li0.05-1.2(PO4)) 407606-26-2. Copper iron lithium phosphate (Cu0-0.8Fe0.2-1Li0.05-1.2(PO4)) 407606-28-4. Aluminum iron lithium phosphate (Al0-0.8Fe0.2-1Li0.05-1.2(PO4)) 407606-30-8. Gallium iron lithium phosphate (Ga0-0.8Fe0.2-1Li0.05-1.2(PO4)) 407606-32-0. Boron iron lithium phosphate (B0-0.8Fe0.2-1Li0.05-1.2(PO4)) 407606-34-2. Iron lithium manganese phosphate (Fe0.2-1Li0.05-1.2Mn0-0.8(PO4)) 407606-36-4. Iron lithium nickel phosphate (Fe0.2-1Li0.05-1.2Ni0-0.8(PO4)) 407606-39-7. Iron lithium vanadium

```
phosphate (Fe0.2-1Li0.05-1.2V0-0.8(PO4)) 407606-42-2, Iron lithium
molybdenum phosphate (Fe0.2-1Li0.05-1.2Mo0-0.8(PO4)) 407606-44-4, Iron
                                                            407606-47-7.
lithium titanium phosphate (Fe0.2-1Li0.05-1.2Ti0-0.8(PO4))
Iron lithium zinc phosphate (Fe0.2-1Li0.05-1.2Zn0-0.8(PO4))
407606-49-9. Iron lithium magnesium phosphate (Fe0.2-1Li0.05-
                  407606-51-3. Iron lithium niobium phosphate
1.2Mq0-0.8(P04)
(Fe0.2-1Li0.05-1.2Nb0-0.8(PO4))
                                 407629-83-8
                                               407629-87-2
                                                             407629-90-7
407629-95-2
             407630-01-7
                           407630-05-1
                                         407630-10-8
                                                       407630-14-2
             407630-25-5. Aluminum iron lithium phosphate
407630-19-7
                     407630-29-9, Gallium iron lithium phosphate
(Al0.7Fe0.3Li(PO4))
(GaO.7FeO.3Li(PO4)) 407630-35-7 407630-40-4. Boron iron
lithium phosphate (B0.75Fe0.25Li(PO4))
                                       407630-46-0
RL: DEV (Device component use); USES (Uses)
   (nonag. electrolyte secondary battery using olivinic lithium phosphorus
   oxide cathode active material)
```

L2 ANSWER 18 OF 30 CAPLUS COPYRIGHT 2004 ACS on STN

ACCESSION NUMBER:

2002:272913 CAPLUS

DOCUMENT NUMBER:

136:297399

TITLE:

Nonaqueous electrolyte secondary battery with a

compound of an olivinic structure as a cathode active

material

INVENTOR(S):

Okawa, Tsuyoshi; Hosoya, Mamoru; Kuyama, Junji;

Fukushima. Yuzuru

PATENT ASSIGNEE(S):

Sony Corporation, Japan Eur. Pat. Appl., 15 pp.

SOURCE:

CODEN: EPXXDW

DOCUMENT TYPE:

Patent

LANGUAGE:

English

FAMILY ACC. NUM. COUNT: 1

PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
	- -			
EP 1195836	A2	20020410	EP 2001-123892	20011005
R: AT, BE, C	CH, DE	, DK, ES, FR	R, GB, GR, IT, LI, LU	, NL, SE, MC, PT,
IE, SI, L	T, LV	, FI, RO		•
JP 2002117833	A2	20020419	JP 2000-308301	20001006
CN 1349266	Α	20020515	CN 2001-142412	20010930
US 2002106564	A1	20020808	US 2001-972395	20011005
US 6656635	B2	20031202		
TW 525312	В	20030321	TW 2001-90124680	20011005
PRIORITY APPLN. INFO.:			JP 2000-308301 A	20001006
AP A non aquoque old	ctrol	uta sacandar	ry cell containing a d	compound of an oli

AB A non-aqueous electrolyte secondary cell containing a compound of an olivinic structure as a cathode active material is to be improved in load characteristics and cell capacity. To this end, there is provided a non-aqueous electrolyte secondary cell including a cathode having a layer of a cathode active material containing a compound represented by the general formula LixFel-yMyPO4, where M is at least one selected from the group consisting of Mn, Cr, Co, Cu, Ni, V, Mo, Ti, Zn, Al, Ga, Mg, B and Nb, with 0.05

ΙT

IT

IT

```
\leq x \leq 1.2 and 0 \leq y \leq 0.8, an anode having a
layer of an anode active material and a non-aqueous electrolyte, wherein the
layer of the cathode active material has a film thickness in a range from
25 to 110 \mu m. If a layer of a cathode active material is provided on
each surface of a cathode current collector, the sum of the film
thicknesses of the layers of the cathode active material ranges between 50
and 220 \mu\text{m}. The non-aqueous electrolyte may be a liquid-based electrolyte or
a polymer electrolyte.
Ball milling
Battery cathodes
Secondary batteries
   (nonag. electrolyte secondary battery with compound of olivinic structure
   as cathode active material)
Carbon black, uses
RL: CPS (Chemical process); DEV (Device component use); PEP (Physical,
engineering or chemical process); PROC (Process); USES (Uses)
   (nonag. electrolyte secondary battery with compound of olivinic structure
   as cathode active material)
10377-52-3. Lithium phosphate 13977-75-8. Phosphoric acid. iron(3+) salt
(3:2)
RL: CPS (Chemical process); PEP (Physical, engineering or chemical
process); PROC (Process)
   (nonag. electrolyte secondary battery with compound of olivinic structure
   as cathode active material)
108-32-7, Propylene carbonate 616-38-6, Dimethyl carbonate
              7782-42-5. Graphite. uses 15365-14-7. Iron lithium
Carbon, uses
phosphate felipo4 21324-40-3. Lithium hexafluorophosphate
                                                              407606-22-8,
Chromium iron lithium phosphate (Cr0-0.8Fe0.2-1Li0.05-1.2(PO4))
407606-24-0, Cobalt iron lithium phosphate (Co0-0.8Fe0.2-1Li0.05-1.2(PO4))
407606-26-2, Copper iron lithium phosphate (Cu0-0.8Fe0.2-1Li0.05-1.2(PO4))
407606-28-4, Aluminum iron lithium phosphate (AlO-0.8Fe0.2-1Li0.05-
           407606-30-8, Gallium iron lithium phosphate
1.2(PO4))
(Ga0-0.8Fe0.2-1Li0.05-1.2(PO4))
                                 407606-32-0. Boron iron lithium
phosphate (B0-0.8Fe0.2-1Li0.05-1.2(P04))
                                           407606-36-4. Iron lithium
nickel phosphate (Fe0.2-1Li0.05-1.2Ni0-0.8(PO4)) 407606-39-7, Iron
lithium vanadium phosphate (Fe0.2-1Li0.05-1.2V0-0.8(PO4))
                                                             407606-42-2.
Iron lithium molybdenum phosphate (Fe0.2-1Li0.05-1.2Mo0-0.8(PO4))
407606-44-4, Iron lithium titanium phosphate (Fe0.2-1Li0.05-1.2Ti0-
            407606-47-7. Iron lithium zinc phosphate (Fe0.2-1Li0.05-1.2Zn0-
0.8(PO4)) 407606-49-9. Iron lithium magnesium phosphate
(Fe0.2-1Li0.05-1.2Mg0-0.8(PO4)) 407606-51-3, Iron lithium niobium
phosphate (Fe'0.2-1Li0.05-1.2Nb0-0.8(PO4))
                                            407629-83-8
                                                           407629-87-2
                            407630-01-7
                                          407630-05-1
                                                        407630-10-8
407629-90-7
              407629-95-2
                            407630-25-5. Aluminum iron lithium phosphate
407630-14-2
              407630-19-7
                      407630-29-9. Gallium iron lithium phosphate
(A10.7Fe0.3Li(P04))
(Ga0.7Fe0.3Li(PO4)) 407630-35-7 407630-40-4. Boron iron
                                         407630-46-0
lithium phosphate (B0.75Fe0.25Li(PO4))
RL: DEV (Device component use); USES (Uses)
   (nonag. electrolyte secondary battery with compound of olivinic structure
   as cathode active material)
```

9011-17-0, Hexafluoropropylene-vinylidene fluoride copolymer RL: DEV (Device component use); MOA (Modifier or additive use); USES (Uses) (nonag. electrolyte secondary battery with compound of olivinic structure as cathode active material) 7439-93-2. Lithium, uses RL: DEV (Device component use); PEP (Physical, engineering or chemical process): PYP (Physical process): PROC (Process): USES (Uses) (nonag. electrolyte secondary battery with compound of olivinic structure as cathode active material) L2 ANSWER 19 OF 30 CAPLUS COPYRIGHT 2004 ACS on STN ACCESSION NUMBER: 2002:272912 CAPLUS DOCUMENT NUMBER: 136:297398 Cathode and anode materials for solid nonaqueous TITLE: electrolyte battery Takahashi, Kimio; Hosoya, Mamoru; Miyake, Masami INVENTOR(S): Sony Corporation, Japan PATENT ASSIGNEE(S): Eur. Pat. Appl., 22 pp. SOURCE: CODEN: EPXXDW DOCUMENT TYPE: Patent English LANGUAGE: FAMILY ACC. NUM. COUNT: 1 PATENT INFORMATION: APPLICATION NO. DATE PATENT NO. KIND DATE _____ 20011004 EP 2001-123773 A2 20020410 EP 1195835 R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT, IE, SI, LT, LV, FI, RO JP 2000-306877 20001005 JP 2002117902 A2 20020419 20020508 CN 2001-138524 20010930 CN 1348231 Α US 2001-969220 20011002 US 2003143465 A1 20030731 TW 522594 В 20030301 TW 2001-90124408 20011003 JP 2000-306877 A 20001005 PRIORITY APPLN. INFO.: A battery is not deteriorated in cell characteristics and maintains the cell shape encapsulated in a laminate film even when overdischarged to a cell voltage of 0 V. The cell includes a cathode containing a compound having the formula LixFe1-yMyPO4, where M is at least one selected from the group consisting of Mn. Cr. Co. Cu. Ni, V. Mo. Ti, Zn. Al, Ga. Mg. B and Nb. with $0.05 \le x \le 1.2$ and $0 \le y \le 0.8$, an anode and a solid electrolyte. A cell member comprised of the cathode and the anode, layered together with the interposition of a solid electrolyte, is encapsulated in a laminate film. Battery anodes IT Battery cathodes Battery electrolytes (cathode and anode materials for solid nonag. electrolyte battery)

7440-44-0. Carbon, uses 15365-14-7. Iron lithium phosphate felipo4 407606-22-8. Chromium iron lithium phosphate (Cr0-0.8Fe0.2-1Li0.05-

TW 513822

CN 1360353

PRIORITY APPLN. INFO.:

В

Α

20021211 20020724

```
1.2(P04))
                 407606-24-0. Cobalt iron lithium phosphate
     (CoO-0.8Fe0.2-1Li0.05-1.2(PO4)) 407606-26-2, Copper iron lithium
     phosphate (Cu0-0.8Fe0.2-1Li0.05-1.2(PO4)) 407606-28-4, Aluminum iron
     lithium phosphate (AlO-0.8Fe0.2-1Li0.05-1.2(PO4))
                                                         407606-30-8, Gallium
     iron lithium phosphate (GaO-0.8FeO.2-1LiO.05-1.2(PO4))
                                                              407606-32-0.
     Boron iron lithium phosphate (B0-0.8Fe0.2-1Li0.05-1.2(PO4))
                                                                   407606-34-2.
     Iron lithium manganese phosphate (Fe0.2-1Li0.05-1.2Mn0-0.8(PO4))
     407606-36-4, Iron lithium nickel phosphate (Fe0.2-1Li0.05-1.2Ni0-0.8(PO4))
     407606-39-7. Iron lithium vanadium phosphate (Fe0.2-1Li0.05-1.2V0-
                407606-42-2. Iron lithium molybdenum phosphate
     0.8(P04)
     (Fe0.2-1Li0.05-1.2Mo0-0.8(PO4))
                                      407606-44-4. Iron lithium titanium
     phosphate (Fe0.2-1Li0.05-1.2Ti0-0.8(PO4)) 407606-47-7, Iron lithium zinc
     phosphate (Fe0.2-1Li0.05-1.2Zn0-0.8(P04)) 407606-49-9. Iron
     lithium magnesium phosphate (Fe0.2-1Li0.05-1.2Mg0-0.8(PO4))
                                                                   407606-51-3.
     Iron lithium niobium phosphate (Fe0.2-1Li0.05-1.2Nb0-0.8(PO4))
     RL: DEV (Device component use); USES (Uses)
        (cathode and anode materials for solid nonaq. electrolyte battery)
     7439-93-2, Lithium, uses
     RL: DEV (Device component use); PEP (Physical, engineering or chemical
     process); PYP (Physical process); PROC (Process); USES (Uses)
        (cathode and anode materials for solid nonag. electrolyte battery)
     ANSWER 20 OF 30 CAPLUS COPYRIGHT 2004 ACS on STN
L2
                         2002:272909 CAPLUS
ACCESSION NUMBER:
DOCUMENT NUMBER:
                         136:297395
TITLE:
                         Method for fabrication of cathode active material and
                         a nonaqueous electrolyte battery
                         Hosoya, Mamoru; Fukushima, Yuzuru; Sakai, Hidecki;
INVENTOR(S):
                         Kuyama, Junji
                         Sony Corporation, Japan
PATENT ASSIGNEE(S):
                         Eur. Pat. Appl., 31 pp.
SOURCE:
                         CODEN: EPXXDW
DOCUMENT TYPE:
                         Patent
LANGUAGE:
                         English
FAMILY ACC. NUM. COUNT: 1
PATENT INFORMATION:
                                           APPLICATION NO. DATE
     PATENT NO.
                      KIND
                           DATE
                                                            20011005
     EP 1195827
                       A2
                            20020410
                                           EP 2001-123894
                            20040310
     EP 1195827
                       Α3
         R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT,
             IE, SI, LT, LV, FI, RO
     JP 2002117848
                       A2
                            20020419
                                           JP 2000-308300
                                                            20001006
                       A2
                            20020419
                                           JP 2000-308313
                                                            20001006
     JP 2002117849
     US 2002124386
                       A1
                            20020912
                                           US 2001-966918
                                                            20010928
```

JP 2000-308300 A 20001006 JP 2000-308313 A 20001006

CN 2001-138169

TW 2001-90124109 20010928

20010930

```
The invention comprises a method for producing a cathode active material
    having superior cell characteristics through single-phase synthesis of a
    composite material composed of a compound represented by the general formula
    LixFel-yMyPO4 and a carbon material pos. and a method for producing a
    non-aqueous electrolyte cell employing the so produced cathode active
    material. To this end, the cathode active material is prepared by a step of
    mixing the starting materials for synthesis of the compound represented by
    the general formula LixFe1-yMyPO4, a step of milling a mixture obtained by
    the mixing step, a step of compressing the mixture obtained by the mixing
    step to a preset d. and a step of sintering the mixture obtained by the
    compressing step. A carbon material is added in any one of the above
    steps prior to the sintering step. The d. of the mixture in the compressing
    step is set to not less than 1.71 g/cm3 and not larger than 2.45 g/cm3.
    Ball milling
ΙT
    Battery cathodes
    Composites
    Secondary batteries
        (method for fabrication of cathode active material and nonaq.
       electrolyte battery)
    Carbon black, uses
IT
    RL: DEV (Device component use); MOA (Modifier or additive use); USES
     (Uses)
        (method for fabrication of cathode active material and nonaq.
        electrolyte battery)
    7440-44-0. Carbon, uses
                              198782-39-7, Iron lithium phosphate
ΙT
     (FeLi0-1(PO4)) 407606-22-8, Chromium iron lithium phosphate
     (Cr0-0.8Fe0.2-1Li0.05-1.2(PO4))
                                     407606-24-0, Cobalt iron lithium
     phosphate (Co0-0.8Fe0.2-1Li0.05-1.2(PO4))
                                                407606-26-2. Copper iron
     lithium phosphate (Cu0-0.8Fe0.2-1Li0.05-1.2(PO4)) 407606-28-4, Aluminum
     iron lithium phosphate (AlO-0.8Fe0.2-1Li0.05-1.2(PO4))
                                                             407606-30-8.
    Gallium iron lithium phosphate (GaO-0.8Fe0.2-1Li0.05-1.2(PO4))
    407606-32-0, Boron iron lithium phosphate (B0-0.8Fe0.2-1Li0.05-1.2(PO4))
    407606-34-2. Iron lithium manganese phosphate (Fe0.2-1Li0.05-1.2Mn0-
                407606-36-4, Iron lithium nickel phosphate
    0.8(P04)
     (Fe0.2-1Li0.05-1.2Ni0-0.8(PO4)) 407606-39-7, Iron lithium vanadium
                                               407606-42-2. Iron lithium
    phosphate (Fe0.2-1Li0.05-1.2V0-0.8(PO4))
    molybdenum phosphate (Fe0.2-1Li0.05-1.2Mo0-0.8(PO4)) 407606-44-4, Iron
    lithium titanium phosphate (Fe0.2-1Li0.05-1.2Ti0-0.8(PO4))
                                                                 407606-47-7.
     Iron lithium zinc phosphate (Fe0.2-1Li0.05-1.2Zn0-0.8(PO4))
     407606-49-9. Iron lithium magnesium phosphate (Fe0.2-1Li0.05-
                       407606-51-3. Iron lithium niobium phosphate
    1.2Mg0-0.8(PO4))
                                      407629-87-2 407629-90-7 407629-95-2
     (Fe0.2-1Li0.05-1.2Nb0-0.8(PO4))
                               407630-14-2
     407630-01-7
                 407630-10-8
    RL: DEV (Device component use); USES (Uses)
        (method for fabrication of cathode active material and nonaq.
        electrolyte battery)
    15365-14-7P. Iron lithium phosphate FeLiPO4
    RL: DEV (Device component use); SPN (Synthetic preparation); PREP
     (Preparation); USES (Uses)
```

(method for fabrication of cathode active material and nonag.

(Uses)

(solid electrolyte cell)

electrolyte battery) 9011-17-0. Hexafluoropropylene-vinylidene fluoride copolymer IT RL: MOA (Modifier or additive use); USES (Uses) (method for fabrication of cathode active material and nonag: electrolyte battery) L2 ANSWER 21 OF 30 CAPLUS COPYRIGHT 2004 ACS on STN ACCESSION NUMBER: 2002:272908 CAPLUS 136:297394 DOCUMENT NUMBER: Solid electrolyte cell TITLE: Goto, Shuji; Hosoya, Mamoru; Endo, Takahiro INVENTOR(S): Sony Corporation, Japan PATENT ASSIGNEE(S): Eur. Pat. Appl., 16 pp. SOURCE: CODEN: EPXXDW DOCUMENT TYPE: Patent English LANGUAGE: FAMILY ACC. NUM. COUNT: 1 PATENT INFORMATION: APPLICATION NO. DATE PATENT NO. KIND DATE A2 20020410 EP 2001-123774 20011004 EP 1195826 Α3 20031126 EP 1195826 R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT, IE, SI, LT, LV, FI, RO JP 2000-306876 20001005 JP 2002117844 A2 20020419 20020718 US 2001-966864 20010928 US 2002094481 A1 TW 2001-90124127 20010928 TW 523952 В 20030311 CN 2001-139323 20010930 CN 1349273 Α 20020515 PRIORITY APPLN. INFO.: JP 2000-306876 A 20001005 A solid electrolyte cell in which cell characteristics are not deteriorated even on overdischarge to the cell voltage of 0 V, such that the shape of the cell encapsulated in the laminate film is maintained. The cell includes a cathode containing a compound represented by the general formula LixFe1-yMyPO4 where $0.05 \le x \le 1.2$, $0 \le y$ ≤ 0.8, and M is at least one selected from the group consisting of Mn, Cr, Co, Cu, Ni, V, Mo, Ti, Zn, Al, Ga, Mg, B and Nb, an anode and a solid electrolyte. An electrode unit 1 comprised of the cathode and the anode layered together with interposition of the solid electrolyte is encapsulated with a laminate film 2. Polyoxyalkylenes, uses IT RL: DEV (Device component use); USES (Uses) (lithium complex; solid electrolyte cell) IT Battery cathodes Secondary batteries (solid electrolyte cell) Fluoropolymers, uses RL: DEV (Device component use); MOA (Modifier or additive use); USES

7439-93-2D, Lithium, polyethylene oxide complex 7791-03-9, Lithium 12031-65-1, Lithium nickel oxide linio2 12057-17-9, perchlorate 15365-14-7. Iron lithium phosphate Lithium manganese oxide limn2o4 25322-68-3D, Polyethylene oxide, lithium complex 116327-69-6. Cobalt lithium nickel oxide CoO.1LiNiO.902 147812-18-8, Iron lithium manganese oxide Fe0.05LiMn1.9504 407606-22-8. Chromium iron lithium phosphate (Cr0-0.8Fe0.2-1Li0.05-1.2(PO4)) 407606-24-0, Cobalt iron lithium phosphate (CoO-0.8Fe0.2-1Li0.05-1.2(PO4)) 407606-26-2. Copper iron lithium phosphate (Cu0-0.8Fe0.2-1Li0.05-1.2(PO4)) 407606-28-4. Aluminum iron lithium phosphate (AlO-0.8Fe0.2-1Li0.05-1.2(PO4)) 407606-30-8, Gallium iron lithium phosphate (GaO-0.8Fe0.2-1Li0.05-407606-32-0, Boron iron lithium phosphate (B0-0.8Fe0.2-1Li0.05-1.2(P04)) 407606-34-2, Iron lithium manganese phosphate 1.2(P04)) 407606-36-4, Iron lithium nickel (Fe0.2-1Li0.05-1.2Mn0-0.8(PO4)) phosphate (Fe0.2-1Li0.05-1.2Ni0-0.8(PO4)) 407606-39-7. Iron lithium 407606-42-2, Iron vanadium phosphate (Fe0.2-1Li0.05-1.2V0-0.8(P04)) lithium molybdenum phosphate (Fe0.2-1Li0.05-1.2Mo0-0.8(PO4)) 407606-44-4, Iron lithium titanium phosphate (Fe0.2-1Li0.05-1.2Ti0-0.8(P04)) 407606-47-7. Iron lithium zinc phosphate (Fe0.2-1Li0.05-1.2Zn0-0.8(PO4)) 407606-49-9. Iron lithium magnesium phosphate (Fe0.2-1Li0.05-1.2Mg0-0.8(PO4)) 407606-51-3. Iron lithium niobium phosphate (Fe0.2-1Li0.05-1.2Nb0-0.8(PO4)) 408331-94-2. Cobalt lithium 408331-95-3. Cobalt lithium manganese nickel oxide ((Co,Ni)Li0-202) oxide ((Co.Mn)Li0-202)408331-96-4. Cobalt lithium zinc oxide 408331-97-5, Cobalt lithium tin oxide ((Co.Sn)Li0-202) ((Co,Zn)Li0-202) 408331-99-7. Cobalt lithium vanadium oxide ((Co,V)Li0-202) 408332-00-3. Cobalt lithium titanium oxide ((Co.Ti)LiO-202) 408332-01-4, Cobalt lithium molybdenum oxide ((Co.Mo)Li0-202) 408332-02-5, Cobalt lithium tungsten oxide ((Co,W)Li0-202) 408332-03-6, Cobalt lithium magnesium 408332-04-7, Cobalt lithium strontium oxide oxide ((Co,Mg)Li0-202) 408332-05-8. Cobalt lithium niobium oxide ((Co.Sr)Li0-202) 408332-06-9. Cobalt iron lithium oxide ((Co,Fe)Li0-202) ((Co,Nb)Li0-202) 408332-07-0, Cobalt copper lithium oxide ((Co,Cu)Li0-202) 408332-08-1. Aluminum cobalt lithium oxide ((Al,Co)Li0-202) 408332-09-2, Cobalt lithium borate oxide (Co0-1Li0-2(B02)0-100-2) 408332-10-5, Cobalt gallium lithium oxide ((Co,Ga)LiO-202) 408332-11-6. Chromium cobalt lithium oxide ((Cr,Co)LiO-202) 408332-12-7, Calcium cobalt lithium oxide 408332-13-8, Iron lithium nickel oxide ((Fe.Ni)Li0-202) ((Ca,Co)Li0-202) 408332-14-9. Copper lithium nickel oxide ((Cu,Ni)Li0-202) 408332-15-0. Aluminum lithium nickel oxide ((Al.Ni)Li0-202) 408332-16-1. Lithium nickel borate oxide (LiO-2NiO-1(BO2)O-100-2) 408332-17-2. Gallium lithium nickel oxide ((Ga,Ni)Li0-202) 408332-18-3, Chromium lithium nickel oxide ((Cr,Ni)Li0-202) 408332-19-4, Calcium lithium nickel oxide ((Ca.Ni)LiO-202) 408332-20-7, Lithium manganese nickel oxide 408332-21-8, Lithium nickel zinc oxide (Li0-2(Ni,Zn)02) (Li0-2(Mn.Ni)02)408332-22-9. Lithium nickel tin oxide (Li0-2(Ni,Sn)02) 408332-23-0. 408332-24-1. Lithium Lithium nickel vanadium oxide (Li0-2(Ni,V)02) nickel titanium oxide (Li0-2(Ni.Ti)02) 408332-25-2, Lithium nickel tungsten oxide (Li0-2(Ni,W)02) 408332-26-3, Lithium molybdenum nickel oxide (LiO-2(Mo.Ni)O2) 408332-27-4, Lithium magnesium nickel oxide

IT

```
408332-28-5. Lithium nickel strontium oxide
     (Li0-2(Mg,Ni)02)
     (Li0-2(Ni,Sr)02)
                        408332-29-6, Lithium nickel niobium oxide
                        408332-30-9. Lithium manganese nickel oxide
     (Li0-2(Ni.Nb)02)
                         408332-31-0. Lithium manganese zinc oxide
     (Li0-2(Mn.Ni)204)
     (Li0-2(Mn,Zn)204)
                         408332-32-1. Lithium manganese tin oxide
                         408332-33-2. Lithium manganese vanadium oxide
     (Li0-2(Mn.Sn)204)
                        408332-34-3. Lithium manganese titanium oxide
     (Li0-2(Mn.V)204)
                         408332-35-4. Lithium manganese molybdenum oxide
     (Li0-2(Mn,Ti)204)
                         408332-36-5, Lithium manganese tungsten oxide
     (Li0-2(Mn.Mo)204)
                        408332-37-6, Lithium magnesium manganese oxide
     (Li0-2(Mn,W)204)
                         408332-38-7. Lithium manganese strontium oxide
     (Li0-2(Mg,Mn)204)
                         408332-39-8, Lithium manganese niobium oxide
     (Li0-2(Mn.Sr)204)
     (Li0-2(Mn, Nb)204)
                         408332-40-1, Iron lithium manganese oxide
                         408332-42-3, Cobalt lithium manganese oxide
     ((Fe,Mn)2Li0-204)
                         408332-44-5. Aluminum lithium manganese oxide
     ((Co.Mn)2Li0-204)
     ((A1.Mn)2Li0-204)
                         408332-45-6. Lithium manganese borate oxide
                                408332-46-7. Gallium lithium manganese oxide
     (Li0-2Mn0-2(B02)0-200-4)
                         408332-47-8, Chromium lithium manganese oxide
     ((Ga,Mn)2Li0-204)
                         408332-48-9. Calcium lithium manganese oxide
     ((Cr.Mn)2Li0-204)
     ((Ca.Mn)2Li0-204)
                         408332-58-1. Aluminum cobalt lithium nickel oxide
                                412351-36-1, Iron lithium manganese phosphate
     (A10.01Co0.98LiNi0.0102)
     (Fe0.9LiMn0.1(PO4))
     RL: DEV (Device component use); USES (Uses)
        (solid electrolyte cell)
                                   108-32-7, Propylene carbonate
                                                                    7782-42-5.
    96-49-1, Ethylene carbonate
                     12190-79-3, Cobalt lithium oxide colio2
                                                                21324-40-3.
     Graphite, uses
     Lithium hexafluorophosphate
                                   24937-79-9. Pvdf
     RL: DEV (Device component use); MOA (Modifier or additive use); USES
     (Uses)
        (solid electrolyte cell)
    ANSWER 22 OF 30 CAPLUS COPYRIGHT 2004 ACS on STN
ACCESSION NUMBER:
                         2002:256757 CAPLUS
DOCUMENT NUMBER:
                         136:282003
                         Lithium-based cathode active materials for
TITLE:
                         rechargeable lithium battery and preparation thereof
INVENTOR(S):
                         Barker, Jeremy; Saidi, M. Yazid; Swoyer, Jeffrey L.
PATENT ASSIGNEE(S):
                         U.S. Pat. Appl. Publ., 39 pp., Cont.-in-part of U.S.
SOURCE:
                         Ser. No. 484.799.
                         CODEN: USXXCO
DOCUMENT TYPE:
                         Patent
LANGUAGE:
                         English
FAMILY ACC. NUM. COUNT:
PATENT INFORMATION:
```

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
US 2002039687	A1	20020404	US 2001-908480	20010718
US 2003129492	A1	20030710	US 2000-484799	20000118

```
WO 2000-US35302 20001222
     WO 2001054212
                      A1
                            20010726
         W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN,
             CR. CU. CZ. DE. DK. DM. DZ. EE, ES, FI. GB. GD. GE. GH. GM. HR.
             HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT,
             LU. LV, MA, MD. MG. MK, MN, MW, MX, MZ, NO, NZ, PL, PT, RO, RU.
             SD, SE, SG, SI, SK, SL, TJ, TM, TR, TT, TZ, UA, UG, US, UZ, VN.
             YU. ZA. ZW. AM. AZ. BY. KG. KZ. MD. RU. TJ. TM
         RW: GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZW, AT, BE, CH, CY,
             DE. DK. ES. FI. FR. GB. GR. IE. IT. LU. MC. NL. PT. SE. TR. BF.
             BJ. CF. CG. CI. CM. GA. GN. GW. ML. MR. NE. SN. TD. TG
                            20030507
                                           EP 2003-2687
                                                            20001222
     EP 1309021
                       A2
                       А3
                            20030903
     EP 1309021
         R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT,
             IE, LT, LV, FI, RO, MK, CY, AL, TR
                                           JP 2002-362497
                                                            20001222
     JP 2003223893
                      A2
                            20030808
PRIORITY APPLN. INFO.:
                                        US 2000-484799
                                                         A2 20000118
                                        WO 2000-US35302 W 20001222
                                        EP 2000-993800
                                                         A3 20001222
                                        JP 2001-553602
                                                         A3 20001222
    The invention provides novel lithium-mixed metal materials which, upon
AB
     electrochem. interaction, release lithium ions, and are capable of
     reversibly cycling lithium ions. The invention provides a rechargeable
     lithium battery which comprises an electrode formed from the novel
     lithium-mixed metal materials. Methods for making the novel lithium-mixed
     metal materials and methods for using such lithium-mixed metal materials
     in electrochem. cells are also provided. The lithium-mixed metal
     materials comprise lithium and at least one other metal besides lithium.
     Preferred materials are lithium-mixed metal phosphates which contain
     lithium and two other metals besides lithium.
ΙT
     Battery cathodes
        (lithium-based cathode active materials for rechargeable lithium
        battery and preparation thereof)
     Olivine-group minerals
IT
     RL: DEV (Device component use); USES (Uses)
        (lithium-based cathode active materials for rechargeable lithium
        battery and preparation thereof)
ΙT
     Secondary batteries
        (lithium; lithium-based cathode active materials for rechargeable
        lithium battery and preparation thereof)
     405914-52-5. Cobalt lithium magnesium phosphate ((Co,Mg)Li(PO4))
IT
     405914-53-6, Cobalt lithium magnesium phosphate (Co0.9LiMg0.1(PO4))
     405914-58-1, Cobalt lithium magnesium phosphate (Co0.95LiMg0.05(PO4))
                  405914-68-3, Calcium cobalt lithium phosphate
     405914-63-8
                        405914-73-0, Calcium cobalt lithium phosphate
     ((Ca,Co)Li(PO4))
     (Ca0.1Co0.9Li(PO4))
                           405914-83-2. Cobalt lithium zinc phosphate
                        405914-88-7. Cobalt lithium zinc phosphate
     ((Co.Zn)Li(PO4))
                           405914-93-4. Cobalt lithium strontium phosphate
     (Co0.9LiZn0.1(PO4))
                        405914-98-9. Cobalt lead lithium phosphate
     ((Co,Sr)Li(PO4))
                        405915-04-0. Cadmium cobalt lithium phosphate
     ((Co,Pb)Li(PO4))
                        405915-09-5. Cobalt lithium tin phosphate
     ((Cd,Co)Li(PO4))
```

```
405915-14-2, Barium cobalt lithium phosphate
        ((Co.Sn)Li(PO4))
                           405915-21-1, Beryllium cobalt lithium phosphate
        ((Ba.Co)Li(PO4))
                           405915-29-9. Cobalt lithium magnesium phosphate
        ((Be.Co)Li(PO4))
                                  405915-34-6, Cobalt lithium magnesium phosphate
        (Co0.5-1LiMq0-0.5(PO4))
                                  405915-39-1, Calcium cobalt lithium phosphate
        (Co0.8-1LiMg0-0.2(P04))
                                  405915-44-8. Calcium cobalt lithium phosphate
        (Ca0-0.5Co0.5-1Li(PO4))
                                  405915-48-2. Cobalt lithium zinc phosphate
        (Ca0-0.2Co0.8-1Li(PO4))
                                  405915-51-7. Cobalt lithium zinc phosphate
        (Co0.5-1LiZn0-0.5(P04))
                                  405915-56-2, Cobalt lithium strontium phosphate
        (Co0.8-1LiZn0-0.2(PO4))
                                  405915-59-5, Cobalt lithium strontium phosphate
        (Co0.5-1LiSr0-0.5(PO4))
                                  405915-63-1. Cobalt lead lithium phosphate
        (Co0.8-1LiSr0-0.2(PO4))
                                  405915-66-4. Cobalt lead lithium phosphate
        (Co0.5-1Pb0-0.5Li(P04))
                                  405915-69-7. Cadmium cobalt lithium phosphate
        (Co0.8-1Pb0-0.2Li(PO4))
                                  405915-79-9. Cadmium cobalt lithium phosphate
        (Cd0-0.5Co0.5-1Li(P04))
                                  405915-82-4, Cobalt lithium tin phosphate
        (Cd0-0.2Co0.8-1Li(P04))
                                  405915-85-7, Cobalt lithium tin phosphate
        (Co0.8-1LiSn0-0.2(PO4))
                                    405915-88-0. Cobalt lithium tin phosphate
        (Co0.95-1LiSn0-0.05(PO4)).
                                  405915-90-4. Barium cobalt lithium phosphate
        (Co0.5-1LiSn0-0.5(PO4))
                                  405915-92-6, Barium cobalt lithium phosphate
        (Ba0-0.5Co0.5-1Li(P04))
                                  405915-94-8. Beryllium cobalt lithium phosphate
        (Ba0-0.2Co0.8-1Li(PO4))
                                  405915-96-0. Beryllium cobalt lithium phosphate
        (Be0-0.5Co0.5-1Li(PO4))
        (Be0-0.2Co0.8-1Li(PO4))
        RL: DEV (Device component use); USES (Uses)
           (lithium-based cathode active materials for rechargeable lithium
           battery and preparation thereof)
        84159-18-2P. Lithium vanadium phosphate Li3V2(PO4)3 349632-76-4P
        . Iron lithium magnesium phosphate (Fe0.9LiMg0.1(PO4))
                                                                  349632-79-7P.
                                                              349632-82-2P, Iron
        Calcium iron lithium phosphate (Ca0.1Fe0.9Li(PO4))
        lithium zinc phosphate (Fe0.9LiZn0.1(PO4))
        RL: DEV (Device component use); SPN (Synthetic preparation); PREP
        (Preparation); USES (Uses)
           (lithium-based cathode active materials for rechargeable lithium
           battery and preparation thereof)
        ANSWER 23 OF 30 CAPLUS COPYRIGHT 2004 ACS on STN
O L2
                            2002:256645 CAPLUS
   ACCESSION NUMBER:
   DOCUMENT NUMBER:
                            136:297382
                            Carbon-coated or carbon-crosslinked redox materials
   TITLE:
                            with transition metal-lithium oxide core for use as
                            battery electrodes
                            Armand, Michel: Gauthier, Michel; Magnan,
   INVENTOR(S):
                            Jean-Francois; Ravet, Nathalie
                            Hydro-Quebec, Can.
   PATENT ASSIGNEE(S):
                            PCT Int. Appl., 78 pp.
   SOURCE:
                            CODEN: PIXXD2
   DOCUMENT TYPE:
                            Patent
                            French
   LANGUAGE:
   FAMILY ACC. NUM. COUNT: 2
   PATENT INFORMATION:
```

```
APPLICATION NO. DATE
     PATENT NO.
                      KIND
                           DATE
                                           -----
                            20020404
                                          WO 2001-CA1350
                                                            20010921
     WO 2002027824
                      A1
        W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN.
            CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, ES, FI, GB, GD, GE, GH,
            GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR.
            LS. LT. LU. LV. MA, MD, MG, MK, MN, MW, MX, MZ, NO, NZ, PH, PL.
            PT. RO. RU. SD. SE, SG. SI, SK, SL, TJ, TM, TR, TT, TZ, UA, UG,
            US. UZ. VN, YU, ZA, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM
         RW: GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZW, AT, BE, CH, CY,
            DE. DK. ES. FI. FR. GB. GR. IE, IT, LU. MC. NL. PT. SE. TR. BF.
             BJ. CF. CG. CI. CM. GA. GN. GQ. GW. ML. MR. NE. SN. TD. TG
                                          AU 2001-93569
     AU 2001093569
                      Α5
                            20020408
                                                            20010921
                                          EP 2001-973907
     EP 1325526
                      Α1
                            20030709
                                                            20010921
        R: AT. BE. CH. DE. DK. ES. FR. GB. GR. IT. LI, LU, NL, SE, MC, PT,
             IE, SI, LT, LV, FI, RO, MK, CY, AL, TR
                                        CA 2000-2320661 A 20000926
PRIORITY APPLN. INFO.:
                                        WO 2001-CA1350
                                                        W 20010921
    Carbon-coated redox materials suitable for use in battery electrodes
AB
     consist of a core surrounded by a coating, or interconnected by carbon
     crosslinks, in which the core includes a composition of formula
     LixM1-yM'y(XO4)n, in which y = 0-0.6, x = 0-2, n = 0-1.5; M is a
     transition metal; and M' is a element of fixed valence selected from Mg2+.
     Ca2+. Al3+. and Zn2+. and X is S, P, and Si. Synthesis of the materials
     is carried out by reacting a balanced mixture of appropriate precursors in a
     reducing atmospheric, to adjust the valence of the transition metals, in the
     presence of a carbon source, which is then pyrolyzed. The resulting
     products exhibit an excellent elec. conductivity and a highly enhanced chemical
     activity.
ΙT
    Silanes
     RL: RCT (Reactant); RACT (Reactant or reagent)
        (alkoxy, silicon source: carbon-coated or carbon-crosslinked redox
       materials with transition metal-lithium oxide core for use as battery
       electrodes)
     Polyoxyalkylenes, uses
ΙT
     RL: NUU (Other use, unclassified); USES (Uses)
        (alkyl ethers, oligomeric, aprotic solvent; carbon-coated or
       carbon-crosslinked redox materials with transition metal-lithium oxide
       core for use as battery electrodes)
IT
     Fluoropolymers, uses
     Polyesters, uses
     Polyethers, uses
     RL: NUU (Other use, unclassified); USES (Uses)
        (binders; carbon-coated or carbon-crosslinked redox materials with
        transition metal-lithium oxide core for use as battery electrodes)
     Battery cathodes
IT
     Battery electrodes
     Redox agents
        (carbon-coated or carbon-crosslinked redox materials with transition
       metal-lithium oxide core for use as battery electrodes)
```

```
Transition metals, uses
IT
     RL: TEM (Technical or engineered material use); USES (Uses)
       (electrodes containing: carbon-coated or carbon-crosslinked redox materials
       with transition metal-lithium oxide core for use as battery electrodes)
    78-93-3. Methyl ethyl ketone, uses 96-48-0. Butyrolactone
ΙT
     Ethylene carbonate 107-21-1D, Ethylene glycol, alkyl ethers 108-32-7.
     Propylene carbonate 111-46-6D. Diethylene glycol, alkyl ethers
     112-27-6D. Triethylene glycol, alkyl ethers 112-60-7D. Tetraethylene
     glycol, alkyl ethers 463-79-6D, Carbonic acid, C1-4-alkyl esters
     RL: NUU (Other use, unclassified); USES (Uses)
        (aprotic solvent; carbon-coated or carbon-crosslinked redox materials
       with transition metal-lithium oxide core for use as battery electrodes)
    9011-14-7. Poly(methyl methacrylate) 24937-79-9. Poly(vinylidene
     difluoride) 25014-41-9. Polyacrylonitrile
     RL: NUU (Other use, unclassified); USES (Uses)
        (binders: carbon-coated or carbon-crosslinked redox materials with
       transition metal-lithium oxide core for use as battery electrodes)
     50-99-7. Glucose, reactions 57-48-7, Fructose, reactions
     Sucrose, reactions 58-86-6, Xylose, reactions 87-79-6, Sorbose
     9002-88-4. Polyethylene 9003-07-0. Polypropylene 9004-34-6. Cellulose.
     reactions 9004-34-6D, Cellulose, esters 9004-35-7, Cellulose acetate
     9005-25-8, Starch, reactions
                                   25212-86-6, Poly(furfuryl alcohol)
     43094-71-9. Ethylene-ethylene oxide copolymer
     RL: RCT (Reactant): RACT (Reactant or reagent)
        (carbon source: carbon-coated or carbon-crosslinked redox materials
       with transition metal-lithium oxide core for use as battery electrodes)
     407640-63-5. Iron lithium titanium phosphate sulfate
IT
     (Fe0.85Li1.35Ti0.15(P04)0.5(S04))
     RL: DEV (Device component use); USES (Uses)
        (electrodes containing; carbon-coated or carbon-crosslinked redox materials
       with transition metal-lithium oxide core for use as battery electrodes)
    7439-89-6D. Iron, mixed oxides 7439-96-5D, Manganese, mixed oxides
IT
     7440-02-0D. Nickel. mixed oxides 7440-32-6D. Titanium, mixed oxides
     7440-47-3D. Chromium, mixed oxides 7440-48-4D. Cobalt, mixed oxides
     7440-50-8D. Copper, mixed oxides 7440-62-2D, Vanadium, mixed oxides
     13816-45-0. Triphylite 15365-14-7, Iron lithium phosphate (FeLiPO4)
     213467-46-0, Iron lithium manganese phosphate (FeLi2Mn(PO4)2)
     RL: TEM (Technical or engineered material use); USES (Uses)
        (electrodes containing; carbon-coated or carbon-crosslinked redox materials
        with transition metal-lithium oxide core for use as battery electrodes)
     90076-65-6
ΙT
     RL: NUU (Other use, unclassified); USES (Uses)
        (electrolyte containing; carbon-coated or carbon-crosslinked redox
        materials with transition metal-lithium oxide core for use as battery
        electrodes)
IT
     516-03-0, Ferrous oxalate
     RL: RCT (Reactant); RACT (Reactant or reagent)
        (iron source: carbon-coated or carbon-crosslinked redox materials with
        transition metal-lithium oxide core for use as battery electrodes)
     7429-90-5, Aluminum, uses 7440-31-5, Tin, uses 7440-36-0, Antimony.
IT
```

IT

IT

IT

IT

IT

reactions

```
7440-66-6, Zinc, uses 7782-42-5, Graphite, uses
                                                          39302-37-9,
Lithium titanate 207803-50-7. Aluminum cobalt lithium magnesium nickel
       258511-24-9. Iron lithium nitride 263898-18-6, Cobalt manganese
nitride · 407640-62-4
RL: DEV (Device component use); USES (Uses)
   (lithium-based cathodes containing; carbon-coated or carbon-crosslinked
  redox materials with transition metal-lithium oxide core for use as
  battery electrodes)
638-38-0, Manganese(II) acetate
RL: RCT (Reactant); RACT (Reactant or reagent)
   (manganese source: carbon-coated or carbon-crosslinked redox materials
  with transition metal-lithium oxide core for use as battery electrodes)
546-89-4. Lithium acetate 553-91-3, Lithium oxalate
                                                       554-13-2, Lithium
                                              1310-65-2. Lithium
carbonate 1309-37-1, Ferric oxide, reactions
          1313-13-9. Manganese dioxide, reactions
                                                    1314-62-1, Vanadium
pentoxide, reactions 1317-61-9. Magnetite, reactions
                                                        10045-86-0.
Ferric phosphate 10102-24-6, Lithium silicate (Li2Si03)
                                                           10377-48-7.
                10377-52-3, Lithium phosphate (Li3PO4)
                                                          10421-48-4,
Lithium sulfate
Ferric nitrate
                12057-24-8. Lithium oxide, reactions
                                                      12627-14-4
13453-80-0. Lithium dihydrogen phosphate 63985-45-5. Lithium
orthosilicate 407640-52-2. Iron lithium manganese phosphate
(Fe0.1-1LiMn0-0.9(PO4)) 407640-53-3, Iron lithium magnesium
phosphate (Fe0.7-1LiMg0-0.3(PO4)) 407640-54-4, Calcium iron lithium
                                   407640-55-5 407640-56-6. Iron
phosphate (Ca0-0.3Fe0.7-1Li(PO4))
lithium phosphate silicate (FeLi1-1.9(PO4)0.1-1(SiO4)0-0.9)
407640-58-8. Iron lithium manganese phosphate sulfate (Fe0-1Li1-1.2Mn0-
0.2[(PO4),(SO4)]) 407640-59-9. Iron lithium manganese phosphate
                      407640-60-2. Iron lithium manganese phosphate
((Fe.Mn)Li1-1.6(P04))
                                        407640-61-3. Iron lithium
sulfate (Fe1-2Li1-2Mn0-1[(P04),(S04)])
titanium phosphate ((Fe,Ti)Li0.5-2(PO4)1.5)
RL: RCT (Reactant): RACT (Reactant or reagent)
   (metal source: carbon-coated or carbon-crosslinked redox materials with
   transition metal-lithium oxide core for use as battery electrodes)
25322-68-3D. Polyethylene glycol, alkyl ethers
RL: NUU (Other use, unclassified); USES (Uses)
   (oligomeric, aprotic solvent; carbon-coated or carbon-crosslinked redox
  materials with transition metal-lithium oxide core for use as battery
   electrodes)
                                       7664-38-2D. Phosphoric acid.
7664-38-2, Phosphoric acid, reactions
       7783-28-0, Ammonium hydrogen phosphate 10124-54-6, Manganese
esters
phosphate
RL: RCT (Reactant); RACT (Reactant or reagent)
   (phosphorus source: carbon-coated or carbon-crosslinked redox materials
   with transition metal-lithium oxide core for use as battery electrodes)
7631-86-9. Silica, reactions
RL: RCT (Reactant); RACT (Reactant or reagent)
   (silicon source: carbon-coated or carbon-crosslinked redox materials
   with transition metal-lithium oxide core for use as battery electrodes)
7664-93-9. Sulfuric acid. reactions 7783-20-2. Ammonium sulfate.
```

RL: RCT (Reactant); RACT (Reactant or reagent)

(sulfur source: carbon-coated or carbon-crosslinked redox materials

with transition metal-lithium oxide core for use as battery electrodes)

REFERENCE COUNT:

THERE ARE 7 CITED REFERENCES AVAILABLE FOR THIS

RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

ANSWER 24 OF 30 CAPLUS COPYRIGHT 2004 ACS on STN

ACCESSION NUMBER:

2002:9172 CAPLUS

DOCUMENT NUMBER:

136:225905

TITLE:

Clustering of Fe3+ in the Li1-3xFexMgPO4 (0<x<0.1)

solid solution

AUTHOR(S):

Goni, Aintzane; Lezama, Luis; Pujana, Ainhoa;

Arriortua, Maria Isabel; Rojo, Teofilo

CORPORATE SOURCE:

Universidad del Pais Vasco, Departamento Quimica

Inorganica, Bilbao, 48080, Spain

SOURCE:

International Journal of Inorganic Materials (2001),

3(7), 937-942

CODEN: IJIMCR: ISSN: 1466-6049

PUBLISHER:

Elsevier Science Ltd.

DOCUMENT TYPE: LANGUAGE:

Journal English

The Li1-3xFexMqP04 (0<x<0.1) solid solution was prepared by solid state synthesis. The structure of these phases was determined by x-ray diffraction on polycryst. samples, being isostructural with LiMgPO4. Fe3+ substitutes part of the Li+ ions in the channels of the LiMgPO4 structure along the [010] direction, creating cation vacancies. The IR bands corresponding to the vibrational modes of the phosphate groups undergo a gradual widening with the amount of inserted iron as a consequence of the increase of disorder in the structure. The EPR spectra show signals with an effective g' = 4.0. This fact can be attributed to the presence of high spin Fe3+ ions in orthorhombic symmetry. The increase of Fe3+ in the compds. leads to a broadening of the Lorentzian EPR signals indicating the existence of magnetic interactions between the Fe3+ ions. Magnetic susceptibility measurements on the Li1-3xFexMgPO4 (0<x<0.1) solid solution show antiferromagnetic behaviors which can be explained considering that the doped Fe3+ ions exhibit a short range magnetic order, forming clusters associated with the vacancies in the structure.

Antiferromagnetic ordering

Crystal structure

ESR (electron spin resonance)

Magnetic susceptibility

Molecular structure

(of lithium iron magnesium phosphate (Li1-3xFexMgPO4 (0<x<0.1)) solid solution containing Fe3+ clusters associated with vacancies)

210709-38-9P, Iron lithium magnesium phosphate (Fe0.03Li0.9MgPO4) ΙT

210709-40-3P, Iron lithium magnesium phosphate (Fe0.1Li0.7MgPO4)

402519-34-0P, Iron lithium magnesium phosphate

(Fe0-0.1Li0.7-1Mg(PO4)) 402519-35-1P, Iron lithium magnesium phosphate (Fe0.07Li0.8Mg(PO4))

RL: PRP (Properties); SPN (Synthetic preparation); PREP (Preparation)

0

JP 2003520182

JP 2003223893

US 2002039687

US 2003077514

PRIORITY APPLN. INFO.:

T2

Α2

A1

A1

20030702

20030808

20020404

20030424

```
(preparation, crystal structure, ESR and magnetic properties)
     1310-65-2. Lithium hydroxide (LiOH) 7722-76-1. Monoammonium phosphate
ΙT
                 10421-48-4. Iron nitrate (Fe(NO3)3)
     RL: RCT (Reactant); RACT (Reactant or reagent)
        (reactant for preparation of lithium iron magnesium phosphate
        (Li1-3xFexMgPO4 (0<x<0.1)) solid solution containing Fe3+ clusters associated
       with vacancies)
                         20
                               THERE ARE 20 CITED REFERENCES AVAILABLE FOR THIS
REFERENCE COUNT:
                               RÊCORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT
    ANSWER 25 OF 30 CAPLUS COPYRIGHT 2004 ACS on STN
L2
                         2001:546025 CAPLUS
ACCESSION NUMBER:
DOCUMENT NUMBER:
                         135:109741
                         Preparation of lithium-based electrochemically active
TITLE:
                         materials for lithium batteries
                         Barker, Jeremy; Saidi, M. Yazid
INVENTOR(S):
                         Valence Technology, Inc., USA
PATENT ASSIGNEE(S):
                         PCT Int. Appl., 97 pp.
SOURCE:
                         CODEN: PIXXD2
DOCUMENT TYPE:
                         Patent
LANGUAGE:
                         English
FAMILY ACC. NUM. COUNT:
                        2
PATENT INFORMATION:
     PATENT NO. .
                                           APPLICATION NO. DATE
                      KIND
                            DATE
                                           -----
                      Α1
                            20010726
                                          WO 2000-US35302 20001222
     WO 2001054212
         W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN.
             CR. CU, CZ, DE, DK, DM, DZ, EE, ES, FI, GB, GD, GE, GH, GM, HR,
             HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT.
             LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NO, NZ, PL, PT, RO, RU,
             SD, SE, SG, SI, SK, SL, TJ, TM, TR, TT, TZ, UA, UG, US, UZ, VN.
             YU, ZA, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM
         RW: GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZW, AT, BE, CH, CY,
             DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, TR, BF,
             BJ, CF, CG, CI, CM, GA, GN, GW, ML, MR, NE, SN, TD, TG
     US 2003129492
                      Α1
                            20030710
                                          US 2000-484799
                                                            20000118
                                           EP 2000-993800
     EP 1252671
                      A1
                            20021030
                                                            20001222
                            20040303
     EP 1252671
                       В1
        R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT,
             IE, SI, LT, LV, FI, RO, MK, CY, AL, TR
                            20030507
                                           EP 2003-2687
                                                            20001222
     EP 1309021
                      A2
     EP 1309021
                      А3
                            20030903
         R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT,
             IE. LT. LV. FI. RO. MK. CY. AL. TR
```

JP 2001-553602

JP 2002-362497

US 2001-908480

US 2002-304890

US 2000-484799

20001222

20001222

20010718

90 20021125 A1 20000118 EP 2000-993800 A3 20001222 JP 2001-553602 A3 20001222 WO 2000-US35302 W 20001222

- AB The invention provides novel lithium-mixed metal materials which, upon electrochem, interaction, release lithium ions, and are capable of reversibly cycling lithium ions. The invention provides a rechargeable lithium battery which comprises an electrode formed from the novel lithium-mixed metal materials. Methods for making the novel lithium-mixed metal materials and methods for using such lithium-mixed metal materials in electrochem, cells are also provided. The lithium-mixed metal materials comprise lithium and at least one other metal besides lithium. Preferred materials are lithium-mixed metal phosphates which contain lithium and two other metals besides lithium.
- IT Secondary batteries

(lithium; preparation of lithium-based electrochem. active materials for lithium batteries)

IT Battery cathodes

(preparation of lithium-based electrochem. active materials for lithium batteries)

- IT Carbon black, uses
 - RL: MOA (Modifier or additive use); USES (Uses)

(preparation of lithium-based electrochem. active materials for lithium batteries)

- IT EPDM rubber
 - RL: TEM (Technical or engineered material use); USES (Uses) (preparation of lithium-based electrochem. active materials for lithium batteries)
- IT 96-49-1, Ethylene carbonate 616-38-6, Dimethyl carbonate 7439-93-2, Lithium, uses 12162-92-4, lithium vanadium oxide liv2o5 21324-40-3, Lithium hexafluorophosphate
 - RL: DEV (Device component use); USES (Uses) (preparation of lithium-based electrochem. active materials for lithium
- IT 331622-65-2P, Iron lithium zinc phosphate (Fe0.8LiZn0.2(PO4))

349632-76-4P, Iron lithium magnesium phosphate (Fe0.9LiMg0.1(P04))

349632-79-7P. Calcium iron lithium phosphate (Ca0.1Fe0.9Li(PO4))

349632-82-2P, Iron lithium zinc phosphate (Fe0.9LiZn0.1(PO4))

349632-85-5P 349632-88-8P

RL: DEV (Device component use); SPN (Synthetic preparation); PREP (Preparation); USES (Uses)

(preparation of lithium-based electrochem. active materials for lithium batteries)

- IT 554-13-2. Lithium carbonate 1305-62-0. Calcium hydroxide. reactions 1309-37-1. Ferric oxide, reactions 1309-42-8. Magnesium hydroxide 7440-44-0. Carbon, reactions 7779-90-0. Zinc phosphate 7783-28-0. Diammonium hydrogen phosphate 10045-86-0. iron phosphate fepo4 13453-80-0. Lithium dihydrogen phosphate 14940-41-1. Iron phosphate fe3(po4)2
 - RL: RCT (Reactant); RACT (Reactant or reagent)
 (preparation of lithium-based electrochem. active materials for lithium

batteries)

IT 15365-14-7P, iron lithium phosphate felipo4 84159-18-2P, Lithium vanadium phosphate Li3V2(PO4)3

RL: SPN (Synthetic preparation); PREP (Preparation)

(preparation of lithium-based electrochem. active materials for lithium batteries)

REFERENCE COUNT:

THERE ARE 16 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

L2 ANSWER 26 OF 30 CAPLUS COPYRIGHT 2004 ACS on STN

ACCESSION NUMBER:

2001:545615 CAPLUS

DOCUMENT NUMBER:

135:109740

TITLE:

Preparation of lithium-containing materials for

battery cathodes

INVENTOR(S):

Barker, Jeremy; Saidi, M. Yazid; Swoyer, Jeffrey L.

PATENT ASSIGNEE(S):

Valence Technology, Inc., USA

SOURCE:

PCT Int. Appl., 94 pp.

CODEN: PIXXD2

DOCUMENT TYPE:

Patent

LANGUAGE:

English

FAMILY ACC. NUM. COUNT:

PATENT INFORMATION:

PATENT NO.		APPLICATION NO. DATE
WO 2001053198	A1 20010726	WO 2000-US35438 20001222
		AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN,
		DZ, EE, ES, FI, GB, GD, GE, GH, GM, HR,
		KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT,
		MN, MW, MX, MZ, NO, NZ, PL, PT, RO, RU,
		TJ. TM. TR, TT, TZ, UA, UG, US, UZ, VN,
		KG, KZ, MD, RU, TJ, TM
		SD, SL, SZ, TZ, UG, ZW, AT, BE, CH, CY,
		GR, IE, IT, LU, MC, NL, PT, SE, TR, BF,
BJ, CF,	CG, CI, CM, GA,	GN, GW, ML, MR, NE, SN, TD, TG
		US 2000-484919 20000118
EP 1252093	A1 20021030	EP 2000-989532 20001222
R: AT, BE,	CH, DE, DK, ES,	FR, GB, GR, IT, LI, LU, NL, SE, MC, PT,
IE, SI,	LT, LV, FI, RO,	MK, CY, AL, TR
		JP 2001-553213 20001222
EP 1391424	A2 20040225	EP 2003-24430 20001222
R: AT, BE,	CH, DE, DK, ES,	FR, GB, GR, IT, LI, LU, NL, SE, MC, PT,
IE, FI.	CY, TR	
US 2002086214	A1 20020704	US 2001-1376 20011019
US 2003215714	A1 20031120	US 2003-461842 20030613
US 6702961	B2 20040309	
PRIORITY APPLN. INFO).:	US 2000-484919 A1 20000118
		EP 2000-989532 A3 20001222
		WO 2000-US35438 W 20001222
		US 2001-1376 A3 20011019

AB The invention provides novel lithium-mixed metal materials which, upon electrochem, interaction, release lithium ions, and are capable of reversibly cycling lithium ions. The invention provides a rechargeable lithium battery which comprises an electrode formed from the novel lithium-mixed metal materials. Methods for making the novel lithium-mixed metal materials and methods for using such lithium-mixed metal materials in electrochem, cells are also provided. The lithium-mixed metal materials comprise lithium and at least one other metal besides lithium. Preferred materials are lithium-mixed metal phosphates which contain lithium and two other metals besides lithium.

IT Reduction

(carbothermic; preparation of lithium-containing materials for battery cathodes)

IT Secondary batteries

(lithium; preparation of lithium-containing materials for battery cathodes)

IT Battery cathodes

(preparation of lithium-containing materials for battery cathodes)

IT 12162-92-4P, lithium vanadium oxide liv2o5 84159-18-2P, Lithium vanadium phosphate Li3V2(P04)3 349632-76-4P, Iron lithium magnesium phosphate (Fe0.9LiMg0.1(P04)) 349632-79-7P. Calcium iron lithium phosphate (Ca0.1Fe0.9Li(P04)) 349632-82-2P, Iron lithium zinc phosphate (Fe0.9LiZn0.1(P04))

RL: DEV (Device component use); SPN (Synthetic preparation); PREP (Preparation); USES (Uses)

(preparation of lithium-containing materials for battery cathodes) 546-89-4, Lithium acetate 553-91-3, Lithium oxalate 554-13-2, Lithium 1305-62-0, Calcium hydroxide, reactions 1309-37-1, Ferric carbonate 1314-62-1, Vanadium 1309-42-8. Magnesium hydroxide oxide, reactions pentoxide. reactions 1317-61-9, iron oxide fe3o4, reactions 7440-44-0. Carbon, reactions 7722-76-1. Ammonium dihydrogen phosphate 7783-28-0. Diammonium hydrogen phosphate 7790-69-4, Lithium nitrate 7803-55-6. Ammonium vanadate 10045-86-0, iron phosphate fepo4 10377-52-3, Lithium 11126-15-1, Lithium vanadium oxide 12036-21-4, Vanadium phosphate 12057-24-8. Lithia, reactions 13453-80-0, Lithium dihydrogen dioxide 15060-59-0. lithium vanadium oxide livo3 phosphate

RL: RCT (Reactant); RACT (Reactant or reagent)

(preparation of lithium-containing materials for battery cathodes)

IT 15365-14-7P, iron lithium phosphate FeLiPO4

RL: SPN (Synthetic preparation), PREP (Preparation)

(preparation of lithium-containing materials for battery cathodes)

REFERENCE COUNT:

THERE ARE 3 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

L2 ANSWER 27 OF 30 CAPLUS COPYRIGHT 2004 ACS on STN

ACCESSION NUMBER:

2001:225610 CAPLUS

DOCUMENT NUMBER:

134:254632

TITLE:

Secondary lithium batteries using lithium iron

phosphate cathodes

INVENTOR(S):

Takahashi, Masaya: Tobishima, Shinichi: Takei, Koji:

Sakurai, Yoji

PATENT ASSIGNEE(S):

Nippon Telegraph and Telephone Corp., Japan

Print selected from Online session 22/03/2004Page 44

SOURCE:

Jpn. Kokai Tokkyo Koho, 8 pp.

CODEN: JKXXAF

DOCUMENT TYPE:

Patent

LANGUAGE:

Japanese

FAMILY ACC. NUM. COUNT: 1

PATENT INFORMATION:

	PATENT NO.	KIND	DATE	А	PPLICATION NO.	. DATE			
		- -		-					
	JP 2001085010			J	P 1999-261394	19990916			
	JP 3504195	В2	20040308						
PRIC	ORITY APPLN. INFO.	:		JP 1	999-261394	19990916			
AB	The batteries us	e LizF	e1-yXyPO4 ((0 < z)	≤ 1; X = eleme	ent .			
	electrochem. sta						ving		
	.olivine-type str								
	is Mg, Co, Ni, a	nd/or	Zn. The ba	atterie	s, capable of	charging and			
	discharging at ≤4 V, inhibit decomposition of electrolyte, and show								
	improved dischar								
IT	Secondary batter		-						
	(secondary Li	batte	ries using	lithiu	m iron phospha	ate cathodes)			
ΙT	Battery cathodes								
	(secondary; s	econda	ry Li batte	eries u	sing lithium [.]	iron phosphate			
	cathodes)								

331622-62-9P, Iron lithium nickel phosphate (Fe0.8LiNi0.2(PO4)) 331622-63-0P, Cobalt iron lithium phosphate (Co0.2Fe0.8Li(PO4))

331622-64-1P, Cobalt iron lithium phosphate (Co0.1Fe0.9Li(PO4))

331622-65-2P, Iron lithium zinc phosphate (Fe0.8LiZn0.2(PO4))

331622-66-3P. Iron lithium magnesium phosphate

(Fe0.85LiMg0.15(PO4))

RL: DEV (Device component use); PNU (Preparation, unclassified); PREP

(Preparation): USES (Uses)

(cathodes; secondary Li batteries using lithium iron phosphate

cathodes)

L2 ANSWER 28 OF 30 CAPLUS COPYRIGHT 2004 ACS on STN

ACCESSION NUMBER:

2001:46145 CAPLUS

DOCUMENT NUMBER:

134:107969

TITLE:

Cleanerless electrophotographic apparatus with

improved durability

INVENTOR(S):

Okado, Kanetsugu; Mizoe, Marekatsu; Arahira, Fumihiro

PATENT ASSIGNEE(S):

Canon Inc., Japan

SOURCE:

Jpn. Kokai Tokkyo Koho, 26 pp.

CODEN: JKXXAF

DOCUMENT TYPE:

Patent Japanese

LANGUAGE:

FAMILY ACC. NUM. COUNT: 1

PATENT INFORMATION:

KIND DATE APPLICATION NO. DATE PATENT NO.

JP 2001013737

A2 20010119 JP 1999-186119

19990630

PRIORITY APPLN. INFO.:

JP 1999-186119

19990630

The invention relates to the cleanerless electrophotog. apparatus which comprises a charging means containing amino-group-containing magnetic particles and showing volume resistivity of 104-109 $\Omega\cdot$ cm, and a development means containing nonmagnetic toner and amino-group-containing spherical magnetic carrier particles. The volume resistivities of the magnetic carrier and the electrophotog. photoconductor are 1010-1015 and $108-1015 \Omega \cdot \text{cm. resp.}$

Electrophotographic apparatus IT

Electrophotographic photoconductors (photoreceptors)

(magnetic particles for cleanerless electrophotog, apparatus with improved durability)

Electrophotographic carriers IT

(magnetic; magnetic particles for cleanerless electrophotog. apparatus with improved durability)

13822-56-5, γ -Aminopropyltrimethoxysilane 26222-39-9. ΙT

Dimethylaminoethyl methacrylate-styrene copolymer

RL: MOA (Modifier or additive use); USES (Uses)

(aminosilane coupling agent on magnetic particle surfaces)

319913-53-6P. Iron lithium magnesium strontium oxide ΙT

RL: PNU (Preparation, unclassified); TEM (Technical or engineered material

use): PREP (Preparation): USES (Uses)

(in magnetic particles for cleanerless electrophotog. apparatus with improved durability)

ANSWER 29 OF 30 CAPLUS COPYRIGHT 2004 ACS on STN L2 \bigcirc

ACCESSION NUMBER:

1998:386203 CAPLUS

DOCUMENT NUMBER:

129:144051

TITLE:

7Li and 31P nuclear magnetic resonance studies of

Li1-3xMgFexP04

AUTHOR(S):

Goni, A.; Bonagamba, T. J.; Silva, M. A.; Panepucci,

H.; Rojo, T.; Barberis, G. E.

CORPORATE SOURCE:

Facultad de Ciencias, Departamento de Quimica

Inorganica, Universidad del Pais Vasco, Bilbao, Spain

SOURCE:

Journal of Applied Physics (1998), 84(1), 416-421

CODEN: JAPIAU: ISSN: 0021-8979

PUBLISHER:

American Institute of Physics

DOCUMENT TYPE:

Journal

English

LANGUAGE:

The authors report a 7Li and 31P NMR study in the Li1-3xMgFexPO4 phases between 150 and 410 K This study, complementary to those made using Moessbauer and magnetic neutron diffraction expts., confirms that the Fe ions enter as Fe(III) in the lattice, and that they enter substituting Li ions. Ionic conductivity measurements, together with the NMR behavior of the 7Li and 31P NMR spectra show that no Li mobility occurs in the temperature range studied even with the addition of the Fe impurity.

Ionic conductivity IT

Mossbauer effect

NMR (nuclear magnetic resonance)

O +

(7Li and 31P NMR studies of Li1-3xMgFexPO4) 7723-14-0, Phosphorus-31, properties 13775-51-4, Lithium magnesium IT phosphate (LiMgPO4) 13982-05-3, Lithium-7, properties 210709-38-9. Iron lithium magnesium phosphate (Fe0.03Li0.9Mg(PO4)) 210709-39-0. Iron lithium magnesium phosphate (Fe0.04Li0.89Mg(PO4)) 210709-40-3, Iron lithium magnesium phosphate (Fe0.1Li0.7Mg(PO4)) RL: PRP (Properties) (7Li and 31P NMR studies of Li1-3xMgFexPO4) THERE ARE 13 CITED REFERENCES AVAILABLE FOR THIS REFERENCE COUNT: 13 RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT L2 ANSWER 30 OF 30 CAPLUS COPYRIGHT 2004 ACS on STN 1993:71061 CAPLUS ACCESSION NUMBER: DOCUMENT NUMBER: 118:71061 Synthesis and conductivity of new lithium-containing TITLE: Nasicon-type phases: Lix[MIIxMIII2-x](SO4)3-y(SeO4)y and Lix[Lix/2MII2-x/2](S04)3-y(Se04)ySlater, Peter R.; Greaves, Colin AUTHOR(S): Sch. Chem., Univ. Birmingham, Birmingham, B15 2TT, UK CORPORATE SOURCE: Journal of Materials Chemistry (1992), 2(12), 1267-9 SOURCE: CODEN: JMACEP; ISSN: 0959-9428 DOCUMENT TYPE: Journal English LANGUAGE: The synthesis of new phases, Lix[MxIIM2-xIII] (SO4)3-y(SeO4)y and Lix[Lix/2M2-x/2III](SO4)3-y(SeO4)y with the Nasicon structure is reported. The ionic conductivities of a range of samples were measured, with the highest conductivity ($\sigma 473 = 1.6 + 10-4 \Omega - 1 \text{ cm} - 1$) being observed for Li0.5[Mg0.5Fe1.5](SO4)3. Attempts to increase the Li content by partially replacing the (S,Se)04 groups by PO4 or SiO4 proved unsuccessful. The synthesis of mixed Na- and Li-containing samples is reported. IT Electric conductors, ceramic (lithium mixed metal sulfates and selenates, preparation and structure of Nasicon-type) Crystal structure IT (of lithium metal sulfates and selenates) IT Electric conductivity and conduction (ionic, of lithium mixed metal sulfates and selenates) 145700-36-3 145700-37-4. Aluminum lithium magnesium sodium ΙT 145481-51-2 sulfate alli0.5mgna0.5(so4)3 145700-38-5, Aluminum lithium magnesium sodium selenate alli0.5mgna0:5(seo4)3 RL: PRP (Properties) (crystal structure and conductivity of) 145481-43-2P, Aluminum lithium nickel sulfate (All.65Li0.35Ni0.35(SO4)3) IT 145481-44-3P. Aluminum lithium zinc sulfate (All.65Li0.35Zn0.35(SO4)3) 145481-46-5P, Aluminum lithium nickel selenate 145481-45-4P 145481-47-6P, Aluminum lithium zinc selenate (All.6Li0.4Ni0.4(Se04)3) (All.6Li0.4Zn0.4(SeO4)3) 145481-48-7P, Iron lithium nickel sulfate (Fel.6Li0.4Ni0.4(SO4)3) 145481-49-8P, Iron lithium zinc sulfate

Print selected from Online session 22/03/2004Page 47

```
145481-50-1P. Chromium lithium nickel sulfate
(Fe1.6Li0.4Zn0.4(SO4)3)
                          145700-27-2P, Aluminum lithium magnesium sulfate
(Cr1.6Li0.4Ni0.4(SO4)3)
                         145700-28-3P. Aluminum lithium sulfate
all.5li0.05mg0.5(so4)3
                  145700-29-4P, Aluminum lithium magnesium selenate
all.8li0.6(so4)3
a11.51i0.5mg0.5(seo4)3
                         145700-30-7P, Aluminum lithium selenate
all.8li0.6(seo4)3 145700-31-8P, Iron lithium magnesium sulfate
                        145700-32-9P. Iron lithium sulfate
fe1.51i0.5mg0.5(so4)3
fe1.81i0.6(so4)3
                  145700-33-0P, Chromium lithium magnesium sulfate
                        145700-34-1P, Chromium lithium zinc sulfate
cr1.51i0.5mg0.5(so4)3
                        145700-35-2P, Chromium lithium sulfate
cr1.51i0.5zn0.5(so4)3
cr1.81i0.6(so4)3
RL: PREP (Preparation)
   (preparation and structure and conductivity of)
```